GenCore version 5.1.6 Copyright (c) 1993 - 2004 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 28, 2004, 20:49:00; Search time 12457 Seconds

(without alignments)

11389.146 Million cell updates/sec

Title: US-10-056-884A-1

Perfect score: 3468

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 2888711 seqs, 20454813386 residues

Total number of hits satisfying chosen parameters: 5777422

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : GenEmbl:*

1: gb_ba:*

2: gb_htg:*

3: gb_in:*

4: gb_om:*

5: gb_ov:*

6: gb_pat:*

7: gb ph:*

8: gb pl:*

9: gb_pr:*

10: gb_ro:*

11: gb sts:*

12: gb_sy:*

13: gb un:*

14: gb vi:*

15: em ba:*

16: em_fun:*

17: em hum:*

18: em_in:*

19: em mu:*

20: em om:*

21: em or:*

22: em ov:*

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26: em_ro:*

27: em_sts:*

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28: em_un:*
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41: em_htgo_other:*
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용

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Res			Query				
	No.	Score	Match	Length	DB	ID	Description
	1	3198.4	92.2	5646	9	AB037738	AB037738 Homo sapi
С	2	2045.4	59.0	182638	2	AC019335	AC019335 Homo sapi
С	3	2043.4	58.9	184589	9	AC008716	AC008716 Homo sapi
С	4	2038.6	58.8	98360	9	AC008473	AC008473 Homo sapi
	5	1640.8	47.3	2412	6	AX405760	AX405760 Sequence
С	6	1161.8	33.5	171949	9	AC008652	AC008652 Homo sapi
	7	1161.8	33.5	209114	9	AC008383	AC008383 Homo sapi
С	8	853.2	24.6	135132	2	AC127249	AC127249 Mus muscu
	9	853.2	24.6	186417	2	AC114984	AC114984 Mus muscu
С	10	819.4	23.6	242679	2	AC117867	AC117867 Rattus no
	11	811.8	23.4	230128	10	AC098707	AC098707 Mus muscu
	12	806.2	23.2	249703	2	AC112599	AC112599 Rattus no
	13	367.4	10.6	781	10	BC049734	BC049734 Mus muscu
	14	319.2	9.2	175059	2	BX323465	BX323465 Danio rer
	15	317.6	9.2	200467	2	BX470157	BX470157 Danio rer
С	16	317.6	9.2	230261	2	BX530085	BX530085 Danio rer
	17	286.4	8.3	184319	2	BX511303	BX511303 Danio rer
	18	284.4	8.2	192400	2	BX530406	BX530406 Danio rer
	19	278.4	8.0	83028	2	BX004755	BX004755 Danio rer
С	20	278.4	8.0	243835	5	AL935304	AL935304 Zebrafish
С	21	255.4	7.4	174712	10	AL831725	AL831725 Mouse DNA
	22	251.6	7.3	1890	5	AY120891	AY120891 Danio rer
	23	220.8	6.4	183038	2	AC107770	AC107770 Mus muscu
С	24	216.4	6.2	246164	2	AC098751	AC098751 Rattus no
	25	216.4	6.2	322972	2	AC129853	AC129853 Rattus no
С	26	212.8		138872	9	AC131951	AC131951 Homo sapi
	27	189.6	5.5	243299	2	AC133800	AC133800 Rattus no
	28	188	5.4	472	6	BD109391	BD109391 EST and e
	29	167	4.8	3086	9	BC013764	BC013764 Homo sapi
С	30	167		109201	9	AC000403	AC000403 Genomic s
С	31	167		169362	2	AL136440	AL136440 Homo sapi
	32	165.4	4.8	251187	2	AL359875	AL359875 Homo sapi
С	33	159.8	4.6	145979	2	AC102815	AC102815 Mus muscu

С	34	156.6	4.5 217336	2 AC131344	AC131344 Rattus no
С	35	149.8	4.3 675	10 BC049679	BC049679 Mus muscu
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	37	128.2	3.7 194240	2 AC118142	AC118142 Rattus no
С	38	127.6	3.7 203720	9 AC093861	AC093861 Homo sapi
С	39	114.8	3.3 569	5 AY093634	AY093634 Acipenser
С	40	104.6	3.0 100029	10 AE014174_3	Continuation (4 of
	41	84.4	2.4 2093	5 BC051776	BC051776 Danio rer
	42	81.8	2.4 1450	17 AF132205	Af132205 Homo sapi
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ALIGNMENTS

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            AB037738
            AB037738.1 GI:7243014
VERSION
KEYWORDS
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  AUTHORS
            Nagase, T., Kikuno, R., Ishikawa, K.I., Hirosawa, M. and Ohara, O.
  TITLE
            Prediction of the coding sequences of unidentified human genes.
            XVI. The complete sequences of 150 new cDNA clones from brain which
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  JOURNAL
            DNA Res. 7 (1), 65-73 (2000)
  MEDLINE
            20181126
   PUBMED
            10718198
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            2
               (bases 1 to 5646)
  AUTHORS
            Ohara, O., Nagase, T. and Kikuno, R.
  TITLE
            Direct Submission
  JOURNAL
            Submitted (31-JAN-2000) Osamu Ohara, Kazusa DNA Research Institute,
            Laboratory of DNA Technology; 1532-3 Yana, Kisarazu, Chiba
            292-0812, Japan (E-mail:cdnainfo@kazusa.or.jp,
            URL:http://www.kazusa.or.jp/huge/, Tel:+81-438-52-3913,
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BASE COUNT ORIGIN 1618 a 1169 c 1150 g 1709 t

Query Match 92.2%; Score 3198.4; DB 9; Length 5646;

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		5; Conservative 0; Mismatches 26; Indels 1; Gaps	1;
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Db	820	AGATTGGATATAGACGAGTTGATTATATTTTATGAAGTAGCAGCTCACTACCATCCACCA	879
Qу	303	TCCAGGGTTTAAACTACTTTTTCAGCATCACTTCACCTGTGGACTCTTATACATTTTGAT	362
Db	880	TCCAGGGTTTAAACTACTTTTTCAGCATCACTTCACCTGTGGACTCTTATACATTTTGAT	939
Qу	363	TTCTTGGGGGAAAATACTGGGATAAGAGGAGGTCATTTTTTAATAAGTTAGCATCCTTT	422
Db	940	TTCTTGGGGGAAAAATACTGGGATAAGAGGAGGTCATTTTTAATAAGTTAGCATCCTTT	999
QУ	423	TCCCTTTCTTACAAGTTGATCCAAAGGATAAGGCTGTGACTCCATTGGATTGCACCTTTA	482
Db	1000	TCCCTTTCTTACAAGTTGATCCAAAGGATAAGGCTGTGACTCCATTGGATTGCACCTTTA	1059
QУ	483	AATCAAAATAGCAGCAGCAGAAGAAAGGGACAATGGCTCTGAGTGGAAACTGTAGTCGTT	542
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Qу	603	ATGTCGGGGGTCAAGTTTATTTTACTCGCCATTCCACATTGATAAGCATCCCTCATTCCC	662
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QУ	783	GGGACAGGCAGGTGGTCCTGCCTGATCACTTTCCAGAAAAAGGAAGACTGAAAAAGGGAAG	842
Db	1360	GGGACAGGCAGGTGCTCCTGATCACTTTCCAGAAAAAGGAAGACTGAAAAAGGGAAG	1419
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Db	1420	CTGAATACTTCCAGCTCCCAGACTTGGTCAAACTCCTGACCCCCGATGAAATCAAGCAAA	1479
Qу	903	GCCCAGATGAATTCTGCCACAGTGACTTTGAAGATGCCTCCCAAGGAAGCGACACAAGAA	962
Db	1480	GCCCAGATGAATTCTGCCACAGTGACTTTGAAGATGCCTCCCAAGGAAGCGACACAAGAA	1539
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Db	1780	TCAAGCACCTGGAAAGGGCTTTTGATATGTTGTCAGAGTGTGGATTCCACATGGTGGCCT	1839
Qу	1263	GTAACTCATCGGTGACAGCATCTTTCATCAACCAATATACAGATGACAAGATCTGGTCAA	1322
Db	1840	GTAACTCATCGGTGACAGCATCTTTCATCAACCAATATACAGATGACAAGATCTGGTCAA	1899
Qy	1323	GCTACACTGAATATGTCTTCTACCGTGAGCCTTCCAGATGGTCACCCTCACACTGCGATT	1382
Db	1900	GCTACACTGAATATGTCTTCTACCGTGAGCCTTCCAGATGGTCACCCTCACACTGCGATT	1959
Qу	1383	GCTGCTGCAAGAATGGCAAAGGTGACAAAGAAGGGGGAGGCGCACGTCTTGCAATGACC	1442
Db		GCTGCTGCAAGAATGGCAAAGGTGACAAAGAAGGGGGAGGCGCACGTCTTGCAATGACC	
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Db	2020	TCTCCACATCTAGCTGCGACAGCCAGTCTGAGGCCAGCTCTCCCCAGGAGACGGTCATCT	2079
Qу		GTGGTCCCGTGACACGCCAGACCAACATCCAGACTCTGGACCGTCCCATCAAGAAGGGCC	
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Qу	1563	CTGTCCAGCTGATCCAACAGTCAGAGATGCGGCGGAAAAGCGACTTACTCCGGATTCTGA	1622
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Db	2320	GGTTTCCTGAGAGAAAACATCCTTGGCAATCTGAACTTTTAAGGAAGTATCATCTATAAG	2379
Qу	1803	GGAGGGCTGGGGGGGAAAAAAAAAAAAAAAGAGTCATTTTGAAATTAACCTCATAAAA	1862
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Qy	1863	${\tt GGAATTCATATTTTAAAGGAAAAAAATACAACTAATGATGCACATTTCTTAGAACACAAT}$	1922
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Qy	1923	AGTCCATTGATATACTACTGCCTACTTTACCTAGTTCACCTTAACATGTAAATCCACAGG	1982
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Qу	2043	CTTCGTCCCATGTGCTAACTATCTTATATATAATGAGAGCCAGCTACGTAAAAGTAGCTG	2102
Db	2619		2678
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Qу	2283	CACTCAAATCTATATGTGCCAGTTTATATTGACTCCGTATGCATGAGTATTTGTGCAACA	2342
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REFERENCE 1 (bases 1 to 182638)
 AUTHORS Waterston, R.H.
 TITLE The sequence of Homo sapiens clone
 JOURNAL Unpublished
REFERENCE 2 (bases 1 to 182638)
 AUTHORS Waterston, R.H.
 TITLE
         Direct Submission
 JOURNAL
          Submitted (01-JAN-2000) Genome Sequencing Center, Washington
          University School of Medicine, 4444 Forest Park Parkway, St. Louis,
         MO 63108, USA
COMMENT
         On Mar 13, 2000 this sequence version replaced gi:6652510.
          ----- Genome Center
          Center: Washington University Genome Sequencing Center
          Center code: WUGSC
          Web site:http://genome.wustl.edu/gsc/index.shtml
          ----- Project Information -----
          Center project name: H NH0427K03
          ----- Summary Statistics -----
          Sequencing vector: M13; 87%
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          Chemistry: Dye-terminator Big Dye; 13% of reads
          Assembly program: Phrap; version 0.990319
          Consensus quality: 174376 bases at least Q40
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          Consensus quality: 178323 bases at least Q20
          Insert size: 182000; agarose-fp
          Insert size: 180938; sum-of-contigs
          Quality coverage: 4.40 in Q20 bases; agarose-fp
          Quality coverage: 4.46 in Q20 bases; sum-of-contigs
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* NOTE: This is a 'working draft' sequence. It currently

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* is not known and their order in this sequence record is
            * arbitrary. Gaps between the contigs are represented as
            * runs of N, but the exact sizes of the gaps are unknown.
            * This record will be updated with the finished sequence
            * as soon as it is available and the accession number will
             be preserved.
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                 41974
                          42073: gap of unknown length
                 42074
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                 50869
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                 50969
                          59100: contig of 8132 bp in length
                 59101
                          59200: gap of unknown length
                 59201
                          72609: contig of 13409 bp in length
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                         126360: contig of 21738 bp in length
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             (bases 1 to 184589)
REFERENCE
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 AUTHORS
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REFERENCE 1 (bases 1 to 98360) AUTHORS DOE Joint Genome Institute and Stanford Human Genome Center.

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 AUTHORS
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          Submitted (03-AUG-1999) Production Sequencing Facility, DOE Joint
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REFERENCE
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 AUTHORS
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COMMENT
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         Tang, Y.T., Liu, C., Zhou, P., Asundi, V., Zhang, J., Zhao, Q.A., Ren, F.,
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         Xue, A.J., Yang, Y., Wehrman, T. and Drmanac, R.T.
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LOCUS AC008652 171949 bp DNA linear PRI 31-JUL-2001

DEFINITION Homo sapiens chromosome 5 clone CTB-18F1, complete sequence.

ACCESSION AC008652

VERSION AC008652.6 GI:15042788

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REFERENCE
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 AUTHORS
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 AUTHORS
          DOE Joint Genome Institute.
 TITLE
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          Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
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* is not known and their order in this sequence record is

* arbitrary. Gaps between the contigs are represented as

* runs of N, but the exact sizes of the gaps are unknown.

* This record will be updated with the finished sequence

* as soon as it is available and the accession number will * be preserved.

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Direct Submission
Submitted (14-MAR-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 186417)
Birren, B., Nusbaum, C., Lander, E., Abouelleil, A., Allen, N.,
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Direct Submission
Submitted (05-JUN-2003) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On May 22, 2003 this sequence version replaced gi:30023906.
All repeats were identified using RepeatMasker:
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http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
    Center: Whitehead Institute/ MIT Center for Genome Research
    Center code: WIBR
    Web site: http://www-seq.wi.mit.edu
    Contact: sequence submissions@genome.wi.mit.edu
----- Project Information
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TITLE

REFERENCE

TITLE

COMMENT

JOURNAL

AUTHORS

JOURNAL

Center project name: L19035 Center clone name: 248 F 9

^{*} NOTE: This is a 'working draft' sequence. It currently

^{*} consists of 6 contigs. The true order of the pieces

^{*} is not known and their order in this sequence record is

^{*} arbitrary. Gaps between the contigs are represented as

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                              11111 11
                                          111111 111 111 111
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DEFINITION
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ACCESSION
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KEYWORDS
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SOURCE
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          Eukarvota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
          Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
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REFERENCE
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  AUTHORS
          Muzny, D. Marie., Metzker, M. Lee., Abramzon, S., Adams, C., Alder, J.,
          Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D.,
          Anyalebechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H.,
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          Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J.,
          Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L.,
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Davila, M.L., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D., Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K., Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K., Egan, A., Escotto, M., Eugene, C., Evans, C.A., Falls, T., Fan, G., Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P., Fraser, C.M., Gabisi, A., Ganta, R., Garcia, A., Garner, T., Garza, M., Gebregeorgis, E., Geer, K., Gill, R., Grady, M., Guerra, W., Guevara, W., Gunaratne, P., Haaland, W., Hamil, C., Hamilton, C., Hamilton, K., Harvey, Y., Havlak, P., Hawes, A., Henderson, N., Hernandez, J., Hernandez, R., Hines, S., Hladun, S.L., Hodgson, A., Hogues, M., Hollins, B., Howells, S., Hulyk, S., Hume, J., Idlebird, D., Jackson, A., Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A., Karpathy, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C., Kowis, C., Kraft, C.L., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J., Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J., Lorensuhewa, L., Loulseged, H., Lozado, R.J., Lu, X., Ma, J., Maheshwari, M., Mahindartne, M., Mahmoud, M., Malloy, K., Mangum, A., Mangum, B., Mapua, P., Martin, K., Martin, R., Martinez, E., Mawhiney, S., McLeod, M.P., McNeill, T.Z., Meenen, E., Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S., Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L., Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S., Nwaokelemeh, O., Okwuonu, G., Olarnpunsagoon, A., Pal, S., Parks, K., Pasternak, S., Paul, H., Perez, A., Perez, L., Pfannkoch, C., Plopper, F., Poindexter, A., Popovic, D., Primus, E., Pu, L.-L., Puazo, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M.A., Reigh, R., Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F., Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S.J., Sanders, W., Savery, G., Scherer, S., Scott, G., Shatsman, S., Shen, H., Shetty, J., Shvartsbeyn, A., Sisson, I., Sitter, C.D., Smajs, D., Sneed, A., Sodergren, E., Song, X.-Z., Sorelle, R., Sosa, J., Steimle, M., Strong, R., Sutton, A., Svatek, A., Tabor, P., Taylor, C., Taylor, T., Thomas, N., Thomas, S., Tingey, A., Trejos, Z., Usmani, K., Valas, R., Vera, V., Villasana, D., Waldron, L., Walker, B., Wang, J., Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F., Williams, G., Willson, R., Wleczyk, R., Wooden, H., Worley, K., Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V., Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von Niederhausern, A., Weiss, R., Smith, D.R., Holt, R.A., Smith, H.O., Weinstock, G. and Gibbs, R.A.

TITLE Direct Submission

JOURNAL Unpublished

REFERENCE 2 (bases 1 to 242679)

AUTHORS Worley, K.C.

TITLE Direct Submission

JOURNAL Submitted (11-APR-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

REFERENCE 3 (bases 1 to 242679)

AUTHORS Rat Genome Sequencing Consortium.

TITLE Direct Submission

JOURNAL Submitted (11-OCT-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

COMMENT On Oct 9, 2002 this sequence version replaced gi:21746224.

The sequence in this assembly is a combination of BAC based reads and whole genome shotgun sequencing reads assembled using Atlas

(http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described in the feature table below represents a scaffold in the Atlas assembly (a 'contig-scaffold'). Within each contig-scaffold, individual sequence contigs are ordered and oriented, and separated by sized gaps filled with Ns to the estimated size. The sequence may extend beyond the ends of the clone and there may be sequence contigs within a contig-scaffold that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.

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shotgun sequence only contigs will be indicated in the feature
table.
----- Genome Center
   Center: Baylor College of Medicine
   Center code: BCM
   Web site: http://www.hgsc.bcm.tmc.edu/
   Contact: hgsc-help@bcm.tmc.edu
----- Project Information
   Center project name: GTZA
   Center clone name: CH230-37619
----- Summary Statistics
   Assembly program: Phrap; version 0.990329
   Consensus quality: 188097 bases at least Q40
   Consensus quality: 190770 bases at least Q30
   Consensus quality: 192614 bases at least Q20
   Estimated insert size: 191086; sum-of-contigs estimation
   Quality coverage: 6x in Q20 bases; sum-of-contigs estimation
* NOTE: Estimated insert size may differ from sequence length
   (see http://www.hgsc.bcm.tmc.edu/docs/Genbank draft data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 11 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
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            204617: contig of 1040 bp in length
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            207199: contig of 1251 bp in length
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            208535: contig of 1236 bp in length
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 Matches 995; Conservative
                             0; Mismatches 166;
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                                                         26; Gaps
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ACCESSION
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VERSION
          AC098707.2 GI:19909459
KEYWORDS
          HTG.
SOURCE
          Mus musculus (house mouse)
 ORGANISM Mus musculus
          Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
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          1 (bases 1 to 230128)
REFERENCE
          McPherson, J.D. and Waterston, R.H.
 AUTHORS
 TITLE
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 JOURNAL
          Unpublished
          2 (bases 1 to 230128)
REFERENCE
          McPherson, J.D. and Waterston, R.H.
 AUTHORS
 TITLE
          Direct Submission
 JOURNAL
          Submitted (31-OCT-2001) Genome Sequencing Center, 4444 Forest Park
          Parkway, St. Louis, MO 63108, USA
REFERENCE
          3 (bases 1 to 230128)
 AUTHORS
          McPherson, J.D. and Waterston, R.H.
 TITLE
          Direct Submission
 JOURNAL
          Submitted (03-APR-2002) Genome Sequencing Center, 4444 Forest Park
          Parkway, St. Louis, MO 63108, USA
REFERENCE
          4 (bases 1 to 230128)
 AUTHORS
          McPherson, J.D. and Waterston, R.H.
 TITLE
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 JOURNAL
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          Parkway, St. Louis, MO 63108, USA
COMMENT
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Contact: submissions@watson.wustl.edu
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ACCESSION AC112599

AC112599.4 GI:23266003 VERSTON

KEYWORDS HTG; HTGS PHASE2; HTGS DRAFT; HTGS_ENRICHED.

SOURCE Rattus norvegicus (Norway rat)

ORGANISM Rattus norvegicus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.

REFERENCE (bases 1 to 249703)

Muzny, D. Marie., Metzker, M. Lee., Abramzon, S., Adams, C., Alder, J., AUTHORS Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D., Anyalebechi, V., Aoyaqi, A., Ayodeji, M., Baca, E., Baden, H., Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F.,

Biswalo, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M., Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E., Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A., Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J., Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L., Davila, M.L., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D., Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K., Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K., Egan, A., Escotto, M., Eugene, C., Evans, C.A., Falls, T., Fan, G.,

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Hollins, B., Howells, S., Hulyk, S., Hume, J., Idlebird, D., Jackson, A., Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A., Karpathy, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C., Kowis, C., Kraft, C.L., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J.,

Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J., Lorensuhewa, L., Loulseged, H., Lozado, R.J., Lu, X., Ma, J.,

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TITLE Direct Submission

JOURNAL Unpublished

(bases 1 to 249703) REFERENCE 2

AUTHORS Worley, K.C.

TITLE Direct Submission

Submitted (22-FEB-2002) Human Genome Sequencing Center, Department **JOURNAL** of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

REFERENCE 3 (bases 1 to 249703)

AUTHORS Rat Genome Sequencing Consortium.

TITLE Direct Submission

JOURNAL Submitted (21-SEP-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One

Baylor Plaza, Houston, TX 77030, USA

On Sep 21, 2002 this sequence version replaced gi:21743383. The sequence in this assembly is a combination of BAC based reads and whole genome shotgun sequening reads assembled using Atlas (http://www.hgsc.bcm.tmc.edu/projects/rat/). As a result, the sequence may extend beyond the ends of the clone and there may be contigs that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.

----- Genome Center

Center: Baylor College of Medicine

Center code: BCM

Web site: http://www.hgsc.bcm.tmc.edu/

Contact: hgsc-help@bcm.tmc.edu

----- Project Information

Center project name: GRQH

Center clone name: CH230-112A20

----- Summary Statistics

Assembly program: Phrap; version 0.990329

Consensus quality: 233268 bases at least Q40

Consensus quality: 235949 bases at least Q30

Consensus quality: 237476 bases at least 020

Estimated insert size: 261159; sum-of-contigs estimation Quality coverage: 4x in Q20 bases; sum-of-contigs estimation

COMMENT

^{*} NOTE: Estimated insert size may differ from sequence length

⁽see http://www.hgsc.bcm.tmc.edu/docs/Genbank draft data.html)

^{*} NOTE: This sequence may represent more than one clone.

^{*} NOTE: This is a 'working draft' sequence. It currently

^{*} consists of 1 contigs. Gaps between the contigs

^{*} are represented as runs of N. The order of the pieces

^{*} is believed to be correct as given, however the sizes

^{*} of the gaps between them are based on estimates that have

^{*} provided by the submittor.

^{*} This sequence will be replaced

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* by the finished sequence as soon as it is available and
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REFEREN AUTHO TITLI JOURN REMAI	ORS S INAL S I	(bases 1 to 781) Strausberg, R. Direct Submission Submitted (31-MAR-2003) National Institutes of Health, Mammalian Gene Collection (MGC), Cancer Genomics Office, National Cancer Enstitute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590, USA WIH-MGC Project URL: http://mgc.nci.nih.gov
COMMENT	r c	Contact: MGC help desk Cmail: cgapbs-r@mail.nih.gov

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cDNA Library Preparation: Michael Brownstein Laboratory
          cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
          DNA Sequencing by: Genome Sequence Centre,
          BC Cancer Agency, Vancouver, BC, Canada
          info@bcgsc.bc.ca
          Steven Jones, Jennifer Asano, Ian Bosdet, Yaron Butterfield,
          Susanna Chan, Readman Chiu, Chris Fjell, Erin Garland, Ran Guin,
          Letticia Hsiao, Martin Krzywinski, Reta Kutsche, Oliver Lee, Soo
          Sen Lee, Victor Ling, Carrie Mathewson, Candice McLeavy, Steven
          Ness, Pawan Pandoh, Anna-Liisa Prabhu, Parvaneh Saeedi, Jacqueline
          Schein, Duane Smailus, Michael Smith, Lorraine Spence, Jeff Stott,
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Qу

Db

Qу

Db

Qy

Db

Qу

Db

Qу

Db

Tissue Procurement: Dr. Jonathan Kuo, NIMH

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coverage: 10.15x in Q20 bases; agarose-fp
          * NOTE: This is a 'working draft' sequence. It currently
          * consists of 3 contigs. The true order of the pieces
          * is not known and their order in this sequence record is
          * arbitrary. Gaps between the contigs are represented as
          * runs of N, but the exact sizes of the gaps are unknown.
          * This record will be updated with the finished sequence
          * as soon as it is available and the accession number will
          * be preserved.
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               44666
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Quality coverage: 10.25x in Q20 bases; sum-of-contigs Quality

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Db 1282	74 GGCGAAAGAAGTTTTCGGCGACGCACTAAACGAGAGCAGGGATCCTGACAGACCGCCGGA
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Db 1283: 128393	34 GCGTTACACTTCTCAGTTTTACCTGAAGTTTCGCCACCTGGAGCGAGC
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Db 1283 128453	94 CGCGGAGAGCGGGTTCCACATCGTCGCGTGCAATTCATCACTCAC
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BX470157.2 GI:30387082
VERSION
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KEYWORDS
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SOURCE
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           1 (bases 1 to 200467)
REFERENCE
 AUTHORS
           McLay, K.
           Direct Submission
 TITLE
           Submitted (04-MAY-2003) Wellcome Trust Sanger Institute, Hinxton,
  JOURNAL
           Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
           zfish-help@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
           On May 5, 2003 this sequence version replaced gi:30349786.
COMMENT
           ----- Genome Center
           Center: Wellcome Trust Sanger Institute
           Center code: SC
           Web site: http://www.sanger.ac.uk
           Contact: zfish-help@sanger.ac.uk
           ----- Project Information
           Center project name: zC119P14
            ----- Summary Statistics
           Assembly program: XGAP4; version 4.5
           Chemistry: Dye-terminator; 100% of reads
           Consensus quality: 198546 bases at least Q40
           Consensus quality: 199010 bases at least Q30
           Consensus quality: 199314 bases at least Q20
           Insert size: 199767; sum-of-contigs
           Insert size: 201190; 3.3% error; agarose-fp
           Quality coverage: 5.95x in Q20 bases; sum-of-contigs Quality
           coverage: 6.07x in Q20 bases; agarose-fp
            _____.
           * NOTE: This is a 'working draft' sequence. It currently
            * consists of 8 contigs. The true order of the pieces
            * is not known and their order in this sequence record is
           * arbitrary. Gaps between the contigs are represented as
            * runs of N, but the exact sizes of the gaps are unknown.
            * This record will be updated with the finished sequence
            * as soon as it is available and the accession number will
            * be preserved.
                          9514: contig of 9514 bp in length
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                          9614: gap of 100 bp
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                         14582: contig of 4968 bp in length
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                14583
                         18933: contig of 4251 bp in length
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                         19033: gap of 100 bp
                18934
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                         66645: contig of 47612 bp in length
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                         66745: gap of 100 bp
                 66746
                         73558: contig of 6813 bp in length
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FEATURES

Location/Qualifiers

1. .200467 source

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Search completed: January 29, 2004, 02:30:04 Job time: 12473 secs

GenCore version 5.1.6 Copyright (c) 1993 - 2004 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 28, 2004, 20:04:04; Search time 867 Seconds

(without alignments)

10797.752 Million cell updates/sec

Title: US-10-056-884A-1

Perfect score: 3468

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

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Total number of hits satisfying chosen parameters: 5105512

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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	8	201	5.8	614	24	ABV95156	Human pancreatic c
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	44	72.6	2.1	346	23	ABV48988	Human prostate exp
	45	72.2	2.1	297	22	AAS29114	cDNA encoding for
	- J	, 2 . 2	۷.1	291	~~	*JUNC 7 T T Z	COMA Elicoding Tot

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XX
AC
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XX
DT
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XX
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XX
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     reproductive disorder; metabolic disorder; premature puberty; nephritis;
KW
KW
     endocrine disorder; memory disorder; neuroendocrine condition; asthma;
KW
     spermatogenesis; renal disease; learning deficiency; Alzheimer's disease;
KW
     neurodegenerative disease; proliferative disorder; autoimmune disease;
KW
     carcinoid tumour; blood coagulation disease; blood platelet disease;
KW
     rheumatoid arthritis; allergy; hyperproliferative disease; gene therapy;
KW
     graft-versus-host disease; organ rejection; antisterility; thrombolytic;
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KW
KW
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DR
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     New potassium channel beta-subunit, K+betaM2, proteins and nucleic
PT
     acids, useful for diagnosing, treating and/or preventing e.g.
PT
     reproductive, neural, metabolic, endocrine, memory, neurodegenerative
PT
     disorders or diseases -
XX
PS
     Claim 1; Page 344-347; 366pp; English.
XX
CC
     The present invention relates to human potassium channel beta-subunit
CC
     (K+betaM2) proteins and polynucleotides encoding such proteins. The
     K+betaM2 sequences are useful for diagnosing, treating and/or preventing
CC
CC
     reproductive disorders, neural disorders, disorders related to aberrant
```

potassium regulation or hyper potassium channel activity, metabolic CC disorders (e.g. premature puberty), endocrine disorders (e.g. aberrant CC growth hormone synthesis and/or secretion), memory disorder, disorders CC of the testis (e.g. spermatogenesis), neuroendocrine condition related CC to aberrant thyroid hormone release, renal disease or disorders (e.g. CC nephritis), disorders related to aberrant higher brain function (e.g. CC learning deficiencies), neurodegenerative diseases (e.g. Alzheimer's CC disease), proliferative disorders (e.g. carcinoid tumour) and disorders CC involving excessive smooth muscle tone or excitability (e.g. asthma). CC CC They may be used to modulate haemostatic or thrombolytic activity, to CC treat or prevent blood coagulation diseases or disorders, blood platelet diseases, wounds, autoimmune diseases, disorders or conditions (e.g. CC CC rheumatoid arthritis), allergic reactions (e.g. asthma), organ rejection CC or graft-versus-host disease, and hyperproliferative diseases. K+betaM2 CC sequences are also used in gene therapy. The present sequence is human CC K+betaM2 cDNA. XX

SQ Sequence 3468 BP; 1038 A; 728 C; 703 G; 999 T; 0 other;

Query Match 100.0%; Score 3468; DB 24; Length 3468; Best Local Similarity 100.0%; Pred. No. 0; Matches 3468; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Qy	121	GATCTGGCAGCTCTGTGTATTTCAGTCAAGTTCCACAATGAAACCTGACAATAATGGTAA	180
Db	121	GATCTGGCAGCTCTGTGTATTTCAGTCAAGTTCCACAATGAAACCTGACAATAATGGTAA	180
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Db	361		420
Qу	421	TTTCCCTTTCTTACAAGTTGATCCAAAGGATAAGGCTGTGACTCCATTGGATTGCACCTT	480
Db	421	TTTCCCTTTCTTACAAGTTGATCCAAAGGATAAGGCTGTGACTCCATTGGATTGCACCTT	480
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Db	481	${\tt TAAATCAAAATAGCAGCAGCAGAAGAAAGGGACAATGGCTCTGAGTGGAAACTGTAGTCG}$	540
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Qу	2281	TTCACTCAAATCTATATGTGCCAGTTTATATTGACTCCGTATGCATGAGTATTTGTGCAA	2340
Db	2281		2340
Qу	2341	CACAAGCACAACTAAGTATGTATATACACATGACGCACACGATGCCAGGGCCTAGACCTC	2400
Db	2341	CACAAGCACAACTAAGTATGTATATACACATGACGCACACGATGCCAGGGCCTAGACCTC	2400
Qу	2401	CCAAGGGCTGTGCTCCTGCTCCCAGCAGCCCTCTCTTAGAATATTTCAGATGGATG	2460
Db	2401		2460
Qу	2461	TCTGACTCTTTCTTAAAATTCTTTTGGGAAGATTTCCCAGCCTTTCTTCACAACACTTTC	2520
Db	2461	TCTGACTCTTTCTTAAAATTCTTTTGGGAAGATTTCCCAGCCTTTCTTCACAACACTTTC	2520
Qу	2521	TAACATCAAATGACTCTCATCATCAACAAATTGTATTCCTTATTGTGAAATTAATACCCT	2580
Db	2521	TAACATCAAATGACTCTCATCATCAACAAATTGTATTCCTTATTGTGAAATTAATACCCT	2580
QУ	2581	CAGGCTCCATTTTACTGCTTTTGCTCTTTGTCTGCATTAAGAGAGGATGAGGAGAGCTGGT	2640
Db	2581	CAGGCTCCATTTTACTGCTCTTTGTCTGCATTAAGAGAGGATGAGGAGAGCTGGT	2640
Qу	2641	CAAACATTCCTTGTGTTAAAAAAATCAAACATTCATATCCACAAAATTTTCTGCTAAATG	2700
Db	2641	CAAACATTCCTTGTGTTAAAAAAATCAAACATTCATATCCACAAAATTTTCTGCTAAATG	2700
QУ	2701	ACTCCACACTCAGCCTTCTCTACCCTGAACTGAATTATCACCCTTTTCTCCATGTTTTCA	2760
Db	2701	ACTCCACACTCAGCCTTCTCTACCCTGAACTGAATTATCACCCTTTTCTCCATGTTTTCA	2760
QУ	2761	GAGTTCTTACTGCCCACAGTTTAATGGTGTGGCCTTTCCACATAATCCACATTAAGTTCT	2820
Db	2761	GAGTTCTTACTGCCCACAGTTTAATGGTGTGGCCTTTCCACATAATCCACATTAAGTTCT	2820
QУ	2821	GTGTTCCTGTGTTGTTGTGGAACTAAGGACAACACACAGTACTTGAATAAGGGTCCGGCC	2880
Db	2821	GTGTTCCTGTGTTGTTGGAACTAAGGACAACACACAGTACTTGAATAAGGGTCCGGCC	2880
Qy	2881	TTTTGTTTGTTTTAGAGAAAGTTGTATTCCACACACACCTAATAATTTCTTATAAAAAT	2940
Db	2881	TTTTGTTTTTAGAGAAAGTTGTATTCCACACACACACATAATTTCTTATAAAAAT	2940
Qу	2941	TTTAAACTACAAAGCTACATTTTTACTTGCTTGTAGCCGTTTTTGTTTG	3000
Db	2941	TTTAAACTACAAAGCTACATTTTTACTTGCTTGTAGCCGTTTTTGTTTG	3000
Qу	3001	CGGGCTTTGGCTGTGCCCATGCTAGGATTTAGCTGTGTCATTTTTATGATGTCTGTAACA	3060
Db	3001	CGGGCTTTGGCTGTGCCCATGCTAGGATTTAGCTGTGTCATTTTTATGATGTCTGTAACA	3060
QУ	3061	ACCCAACAAGGTAACTGAAGCTCCAGAGTTAAGGTTTCAGATTTCTAAATGAAACTATCT	3120

```
3061 ACCCAACAAGGTAACTGAAGCTCCAGAGTTAAGGTTTCAGATTTCTAAATGAAACTATCT 3120
Db
      3121 TTTTCAATTACATCCTGACTTGTATAGACACAGCCAAAAAGAAACTGTTAATAGCCATCC 3180
Qу
         3121 TTTTCAATTACATCCTGACTTGTATAGACACAGCCAAAAAGAAACTGTTAATAGCCATCC 3180
Db
      3181 GTCCATGTAACTCTGTATTTTACTAAGGTACCAATAGCTCTTTCATAGACTTGTGCTACA 3240
QУ
         Db
      3181 GTCCATGTAACTCTGTATTTTACTAAGGTACCAATAGCTCTTTCATAGACTTGTGCTACA 3240
      3241 AGAAGGTTAAAAGACCAGTTTTATTTTCAGCATTCCTCATGCATTTCAGTGGTAACCAAA 3300
Qу
         Db
      3241 AGAAGGTTAAAAGACCAGTTTTATTTTCAGCATTCCTCATGCATTTCAGTGGTAACCAAA 3300
      3301 AAATAATTTGTCAATTAATAGTTGTGTGCCAAGCACTCCTAATTTGTTTTATTGCGTGTG 3360
Qу
         Db
      3301 AAATAATTTGTCAATTAATAGTTGTGTGCCAAGCACTCCTAATTTGTTTTATTGCGTGTG 3360
      Qу
         Db
Qу
      Db
RESULT 2
ABN59764
ID
   ABN59764 standard; cDNA; 2412 BP.
XX
AC
   ABN59764;
XX
DT
   28-JUN-2002 (first entry)
XX
DE
   Novel human coding sequence SEQ ID NO: 175.
XX
KW
   Human; antianaemic; vulnerary; antiinflammatory; immunomodulator;
KW
   antiinfertility; cerebroprotective; cytostatic; rheumatic; gene therapy;
KW
   neuroprotective; antiparkinsonian; protein therapy; EST;
KW
   expressed sequence tag; gene; ss.
XX
os
   Homo sapiens.
XX
   WO200222660-A2.
PN
XX
PD
   21-MAR-2002.
XX
PF
   10-SEP-2001; 2001WO-US26015.
XX
   11-SEP-2000; 2000US-0659671.
PR
XX
PΑ
   (HYSE-) HYSEQ INC.
XX
PΙ
   Tang YT, Liu C, Zhou P, Asundi V, Zhang J,
                                    Zhao QA,
                                         Ren F;
PΙ
   Xue AJ, Yang Y, Wehrman T, Drmanac RT;
```

```
XX
    WPI: 2002-292408/33.
DR
    P-PSDB; ABB97351.
DR
XX
    An isolated polynucleotide for treating diseases associated with its
PT
    encoded polypeptide such as cancer and multiple sclerosis -
PT
XX
    Claim 1; SEQ ID NO 175; 509pp; English.
PS
XX
    The present invention provides the protein and coding sequences of 444
CC
    novel human proteins. These were isolated from expressed sequences tags
CC
    (ESTs). They can be used to stimulate cell growth, to regulate
CC
CC
    haematopoiesis e.g. to treat aplastic anaemia, to help tissue regrowth
CC
    e.g. in burn treatment, to regulate the immune system e.g. to treat
    multiple sclerosis, to regulate activin or inhibin e.g. to treat
CC
CC
    infertility, to regulate haemostasis or thrombolysis e.g. to treat
CC
    stroke and cancer, to screen for drugs, to treat inflammatory conditions
    e.g. rheumatoid arthritis, and to treat nervous system disorders e.g.
CC
CC
    Parkinson's disease. The present sequence is a coding sequence of the
CC
    invention.
XX
    Sequence 2412 BP; 638 A; 585 C; 551 G; 638 T; 0 other;
SO
 Query Match
                     47.3%;
                           Score 1640.8; DB 24; Length 2412;
 Best Local Similarity
                    99.6%;
                           Pred. No. 0:
 Matches 1645; Conservative
                          0: Mismatches
                                         7: Indels
                                                    0;
                                                       Gaps
                                                             0;
Qy
        Db
        243 AGATTGGATATAGACGAGTTGATTATATTTTATGAAGTAGCAGCTCACTACCATCCACCA 302
Qу
           820 AGATTGGATATAGACGAGTTGATTATATTTTATGAAGTAGCAGCTCACTACCATCCACCA 879
Db
        303 TCCAGGGTTTAAACTACTTTTTCAGCATCACTTCACCTGTGGACTCTTATACATTTTGAT 362
Qy
           880 TCCAGGGTTTAAACTACTTTTTCAGCATCACTTCACCTGTGGACTCTTATACATTTTGAT 939
Db
Qy
        363 TTCTTGGGGGAAAAATACTGGGATAAGAGGAGGTCATTTTTTAATAAGTTAGCATCCTTT 422
           940 TTCTTGGGGGAAAATACTGGGATAAGAGGAGGTCATTTTTTAATAAGTTAGCATCCTTT 999
Db
Qу
        423 TCCCTTTCTTACAAGTTGATCCAAAGGATAAGGCTGTGACTCCATTGGATTGCACCTTTA 482
           1000 TCCCTTTCTTACAAGTTGATCCAAAGGATAAGGCTGTGACTCCATTGGATTGCACCTTTA 1059
Db
Qу
        483 AATCAAAATAGCAGCAGCAGAAGAAAGGGACAATGGCTCTGAGTGGAAACTGTAGTCGTT 542
           1060 AATCAAAATAGCAGCAGCAGAAGAAAGGGACAATGGCTCTGAGTGGAAACTGTAGTCGTT 1119
Db
Qу
        543 ATTATCCTCGAGAACAAGGGTCCGCAGTTCCCAACTCCTTCCCTGAGGTGGTAGAGCTGA 602
           1120 ATTATCCTCGAGAACAAGGGTCCGCAGTTCCCAACTCCTTCCCTGAGGTGGTAGAGCTGA 1179
Db
        603 ATGTCGGGGGTCAAGTTTATTTTACTCGCCATTCCACATTGATAAGCATCCCTCATTCCC 662
Qу
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Db	1180	ATGTCGGGGGTCAAGTTTATTTTACTCGCCATTCCACATTGATAAGCATCCCTCATTCCC	1239
Qу	663	TCCTGTGGAAAATGTTTTCCCCAAAGAGAGACACGGCTAATGATCTAGCCAAGGACTCCA	722
Db	1240	TCCTGTGGAAAATGTTTTCCCCAAAGAGACACGGCTAATGATCTAGCCAAGGACTCCA	1299
Qу	723	AGGGAAGGTTTTCATTGACAGAGATGGATTCTTGTTCCGTTATATTCTGGACTATCTCA	782
Db	1300	AGGGAAGGTTTTTCATTGACAGAGATGGATTCTTGTTCCGTTATATTCTGGACTATCTCA	1359
Qу	783	GGGACAGGCAGGTGGTCCTGCCTGATCACTTTCCAGAAAAAGGAAGACTGAAAAGGGAAG	842
Db	1360	GGGACAGGCAGGTGGTCCTGATCACTTTCCAGAAAAAGGAAGACTGAAAAAGGGAAG	1419
Qу	843	CTGAATACTTCCAGCTCCCAGACTTGGTCAAACTCCTGACCCCCGATGAAATCAAGCAAA	902
Db	1420	CTGAATACTTCCAGCTCCCAGACTTGGTCAAACTCCTGACCCCCGATGAAATCAAGCAAA	1479
Qy	903	GCCCAGATGAATTCTGCCACAGTGACTTTGAAGATGCCTCCCAAGGAAGCGACACAAGAA	962
Db	1480	GCCCAGATGAATTCTGCCACAGTGACTTTGAAGATGCCTCCCAAGGAAGCGACACAAGAA	1539
Qу	963	TCTGCCCCCTTCCTCCTGCCGCCGCCGCAGTGGGGTTTCATTACTGTGGGTT	1022
Db	1540	TCTGCCCCCTTCCTCCCTGCCCGACCGCAAGTGGGGTTTCATTACTGTGGGTT	1599
Qy	1023	ACAGAGGATCCTGCACCTTGGGCAGAGGGGACAGGCAGATGCCAAGTTTCGGAGAGTTC	1082
Db	1600		1659
QУ	1083	CCCGGATTTTGGTTTGTGGAAGGATTTCCTTGGCAAAAGAAGTCTTTGGAGAAACTTTGA	1142
Db	1660	CCCGGATTTTGGTGGAAGGATTTCCTTGGCAAAAGAAGTCTTTGGAGAAACTTTGA	1719
Qу	1143	ATGAAAGCAGAGACCCTGATCGAGCCCCAGAAAGATACACCTCCAGATTTTATCTCAAAT	1202
Db	1720	ATGAAAGCAGAGACCCTGATCGAGCCCCAGAAAGATACACCTCCAGATTTTATCTCAAAT	1779
Qу	1203	TCAAGCACCTGGAAAGGGCTTTTGATATGTTGTCAGAGTGTGGATTCCACATGGTGGCCT	1262
Db	1780	TCAAGCACCTGGAAAGGGCTTTTGATATGTTGTCAGAGTGTGGATTCCACATGGTGGCCT	1839
Qу	1263	GTAACTCATCGGTGACAGCATCTTTCATCAACCAATATACAGATGACAAGATCTGGTCAA	1322
Db	1840	GTAACTCATCGGTGACAGCATCTTTCATCAACCAATATACAGATGACAAGATCTGGTCAA	1899
Qу	1323	GCTACACTGAATATGTCTTCTACCGTGAGCCTTCCAGATGGTCACCCTCACACTGCGATT	1382
Db	1900	GCTACACTGAATATGTCTTCTACCGTGAGCCTTCCAGATGGTCACCCTCACACTGCGATT	1959
Qу	1383	GCTGCTGCAAGAATGGCAAAGGTGACAAAGAAGGGGGAGGGGCACGTCTTGCAATGACC	1442
Db	1960	GCTGCTGCAAGAATGGCAAAGGTGACAAAGAAGGGGGAGAGCGGCACGTCTTGCAATGACC	2019
Qу	1443	TCTCCACATCTAGCTGCGACAGCCAGTCTGAGGCCAGCTCTCCCCAGGAGACGGTCATCT	1502
Db	2020	TCTCCACATCTAGCTGCGACAGCCAGTCTGAGGCCAGCTCTCCCCAGGAGACGGTCATCT	2079

```
1503 GTGGTCCCGTGACACGCCAGACCAACATCCAGACTCTGGACCGTCCCATCAAGAAGGGCC 1562
Qу
          2080 GTGGTCCCGTGACACGCCAGACCAACATCCAGACTCTGGACCGTCCCATCAAGAAGGGCC 2139
Db
Qу
       1563 CTGTCCAGCTGATCCAACAGTCAGAGATGCGGCGGAAAAGCGACTTACTCCGGATTCTGA 1622
          2140 CTGTCCAGCTGATCCAACAGTCAGAGATGCGGCGGAAAAGCGACTTACTCCGGACTCTGA 2199
Db
       Qу
          Db
       Qу
          2260 CAATTGAGGAGGAGCTGGAGAAATGTATCCAGGATTTCCTAAAAATCAAAATTCCAGATC 2319
Db
       1743 GGTTTCCTGAGAGAAACATCCTTGGCAATCTGAACTTTTAAGGAAGTATCATCTATAAG 1802
Qу
          Db
       2320 GGTTTCCTGAGAGAAAACATCCTTGGCAATCTGAACTTTTAAGGAAGTATCATCTATAAG 2379
       1803 GGAGGGCTGGGGGGGGGAAAAAAAAAAAAAA 1834
Qy
           2380 GGAGGGCTGGGGGGGGAAAAGAAAAAAAA 2411
Dh
RESULT 3
AAD46125
ID
   AAD46125 standard; DNA; 769 BP.
XX
AC
   AAD46125;
XX
DT
   27-DEC-2002 (first entry)
XX
DE
   Human BAC AC008652 exon used to isolate K+betaM2 cDNA.
XX
   Human; potassium channel beta-subunit; K+betaM2 protein; neural disorder;
KW
KW
    reproductive disorder; metabolic disorder; premature puberty; nephritis;
   endocrine disorder; memory disorder; neuroendocrine condition; asthma;
KW
KW
    spermatogenesis; renal disease; learning deficiency; Alzheimer's disease;
   neurodegenerative disease; proliferative disorder; autoimmune disease;
KW
    carcinoid tumour; blood coagulation disease; blood platelet disease;
KW
KW
    rheumatoid arthritis; allergy; hyperproliferative disease; gene therapy;
KW
    graft-versus-host disease; organ rejection; antisterility; thrombolytic;
KW
   antiinflammatory; neuroprotective; anti-Parkinsonian; immunosuppressive;
KW
   nephrotropic; cytostatic; nootropic; hypotensive; vulnerary; ds.
XX
   Homo sapiens.
OS
XX
PN
   WO200266601-A2.
XX
PD
   29-AUG-2002.
XX
PF
    24-JAN-2002; 2002WO-US02332.
XX
PR
   24-JAN-2001; 2001US-263872P.
PR
    14-FEB-2001; 2001US-269794P.
```

```
XX
PA
    (BRIM ) BRISTOL-MYERS SQUIBB CO.
XX
PΙ
             Lee L, Chen J, Jackson D, Ramanathan C, Siemers N;
PΙ
    Chang H,
             Carroll P:
XX
    WPI; 2002-691617/74.
DR
XX
РТ
    New potassium channel beta-subunit, K+betaM2, proteins and nucleic
PT
    acids, useful for diagnosing, treating and/or preventing e.g.
PТ
    reproductive, neural, metabolic, endocrine, memory, neurodegenerative
    disorders or diseases -
PT
XX
PS
    Example 1; Page 349-350; 366pp; English.
XX
CC
    The present invention relates to human potassium channel beta-subunit
CC
    (K+betaM2) proteins and polynucleotides encoding such proteins. The
    K+betaM2 sequences are useful for diagnosing, treating and/or preventing
CC
CC
    reproductive disorders, neural disorders, disorders related to aberrant
CC
    potassium regulation or hyper potassium channel activity, metabolic
CC
    disorders (e.g. premature puberty), endocrine disorders (e.g. aberrant
CC
    growth hormone synthesis and/or secretion), memory disorder, disorders
CC
    of the testis (e.g. spermatogenesis), neuroendocrine condition related
CC
    to aberrant thyroid hormone release, renal disease or disorders (e.g.
CC
    nephritis), disorders related to aberrant higher brain function (e.g.
CC
    learning deficiencies), neurodegenerative diseases (e.g. Alzheimer's
CC
    disease), proliferative disorders (e.g. carcinoid tumour) and disorders
CC
    involving excessive smooth muscle tone or excitability (e.g. asthma).
CC
    They may be used to modulate haemostatic or thrombolytic activity, to
    treat or prevent blood coagulation diseases or disorders, blood platelet
CC
CC
    diseases, wounds, autoimmune diseases, disorders or conditions (e.g.
CC
    rheumatoid arthritis), allergic reactions (e.g. asthma), organ rejection
CC
    or graft-versus-host disease, and hyperproliferative diseases. K+betaM2
CC
    sequences are also used in gene therapy. The present sequence is human
CC
    BAC AC008652 exon used to isolate K+betaM2 cDNA. This sequence is used
CC
    in the exemplification of the invention.
XX
SO
    Sequence 769 BP; 209 A; 180 C; 184 G; 196 T; 0 other;
 Query Match
                        22.2%; Score 769; DB 24; Length 769;
 Best Local Similarity
                       100.0%; Pred. No. 6.5e-143;
 Matches 769; Conservative
                             0; Mismatches
                                               0; Indels
                                                           0; Gaps
                                                                       0;
         393 AGGTCATTTTTTAATAAGTTAGCATCCTTTTCCCTTTCTTACAAGTTGATCCAAAGGATA 452
Qу
             Db
           1 AGGTCATTTTTAATAAGTTAGCATCCTTTTCCCTTTCTTACAAGTTGATCCAAAGGATA 60
         453 AGGCTGTGACTCCATTGGATTGCACCTTTAAATCAAAATAGCAGCAGCAGAAGAAAGGGA 512
Qу
             Db
          61 AGGCTGTGACTCCATTGGATTGCACCTTTAAATCAAAATAGCAGCAGCAGAAGAAAGGGA 120
         513 CAATGGCTCTGAGTGGAAACTGTAGTCGTTATTATCCTCGAGAACAAGGGTCCGCAGTTC 572
Qу
             Db
         121 CAATGGCTCTGAGTGGAAACTGTAGTCGTTATTATCCTCGAGAACAAGGGTCCGCAGTTC 180
         573 CCAACTCCTTCCCTGAGGTGGTAGAGCTGAATGTCGGGGGTCAAGTTTATTTTACTCGCC 632
Qу
```

Db	181	${\tt CCAACTCCTTCCCTGAGGTGGTAGAGCTGAATGTCGGGGGTCAAGTTTATTTTACTCGCC}$	240					
Qу	633	ATTCCACATTGATAAGCATCCCTCATTCCCTCTGTGGAAAATGTTTTCCCCAAAGAGAG	692					
Db	241		300					
Qу	693		752					
Db	301	ACACGGCTAATGATCTAGCCAAGGACTCCAAGGGAAGGTTTTTCATTGACAGAGATGGAT	360					
Qу	753	TCTTGTTCCGTTATATTCTGGACTATCTCAGGGACAGGCAGG	812					
Db	361	TCTTGTTCCGTTATATTCTGGACTATCTCAGGGACAGGCAGG	420					
Qу	813	TTCCAGAAAAAGGAAGACTGAAAAGGGAAGCTGAATACTTCCAGCTCCCAGACTTGGTCA	872					
Db	421	TTCCAGAAAAGGAAGACTGAAAAGGGAAGCTGAATACTTCCAGCTCCCAGACTTGGTCA	480					
Qу	873	AACTCCTGACCCCGATGAAATCAAGCAAAGCCCAGATGAATTCTGCCACAGTGACTTTG	932					
Db	481		540					
Qу	933	AAGATGCCTCCCAAGGAAGCGACACAAGAATCTGCCCCCCTTCCTCCCTGCCG	992					
Db	541	AAGATGCCTCCCAAGGAAGCGACACAAGAATCTGCCCCCCTTCCTCCCTGCTCCCTGCCG	600					
Qу	993	ACCGCAAGTGGGGTTTCATTACTGTGGGTTACAGAGGATCCTGCACCTTGGGCAGAGAGG	1052					
Db	601	ACCGCAAGTGGGGTTTCATTACTGTGGGTTACAGAGGATCCTGCACCTTGGGCAGAGAG	660					
Qу	1053	GACAGGCAGATGCCAAGTTTCGGAGAGTTCCCCGGATTTTGGTTTGTGGAAGGATTTCCT	1112					
Db	661		720					
Qу	1113	TGGCAAAAGAAGTCTTTGGAGAAACTTTGAATGAAAGCAGAGACCCTGA 1161						
Db	721							
RESU								
ID	9216/c ABA09216	standard; cDNA; 906 BP.						
XX	. 10.10.52.10	Sanata, Opini, 300 DL.						
AC	ABA09216;							
XX	XX DT 11-JAN-2002 (first entry)							
XX								

```
DE
     Human VM106R.1 homologue-encoding cDNA, SEQ ID NO:992.
XX
KW
     Human; cytokine; cell proliferation; cell differentiation; growth factor;
KW
     haematopoiesis regulation; tissue growth; immunomodulator; activin;
KW
     inhibin; chemotaxis; chemokinesis; thrombolysis; oncogenesis;
     proliferation; metastasis; cancer; tumour; haematopoietic disorder;
KW
KW
    myeloid cell disorder; lymphoid cell disorder; asthma; arthritis;
KW
     chronic inflammatory condition; proliferative retinopathy;
KW
     atherosclerosis; coronary heart disease; arterial ischaemia;
KW
     bone disorder; osteoporosis; vascular growth disorder;
```

KW tissue regeneration; wound healing; infection; immune disorder; cell culture; drug screening; gene therapy; antiinflammatory; ΚW antiasthmatic; antiarthritic; haemostatic; antiarteriosclerotic; KW KW cytostatic; osteopathic; vasotropic; cardiant; virucide; antibacterial; KW antifungal; vulnerary; antiulcer; ss. XX os Homo sapiens. XX PN WO200157188-A2. XX PD 09-AUG-2001. XX PF05-FEB-2001; 2001WO-US03800. XX PR 03-FEB-2000; 2000US-0496914. 27-APR-2000; 2000US-0560875. PR XX PA (HYSE-) HYSEQ INC. XX PΙ Tang YT, Liu C, Drmanac RT; XX DR WPI; 2001-457740/49. DR P-PSDB; ABB11972. XX PTHuman proteins and DNA encoding sequences useful for preventing, PTtreating or ameliorating a medical condition in a mammalian subject PTe.g. arthritis and cancer -XX PS Claim 1; Page 844-845; 1963pp; English. XX CC Sequences ABB10981-ABB12330 represent 1350 novel human polypeptides, and CC sequences ABA08225-ABA09574 represent nucleic acids encoding them. The invention also relates to vectors and recombinant host cells comprising a CC CC nucleotide of the invention, methods of producing the novel polypeptides, CC antibodies against the polypeptides, methods of detecting the nucleotides CC or polypeptides in a sample, and methods of identifying compounds which CC bind to polypeptides of the invention. Although novel, many of the CC polypeptides of the invention have homology to known proteins, thereby CC giving an insight into their probable biological activities, and hence CC potential therapeutic applications. The polypeptides of the invention may CC have various activities, including cytokine, cell proliferation or cell differentiation activities; stem cell growth factor activity; CC haematopoiesis regulatory activity; tissue growth activity; CC CC immunomodulatory activity; activin- or inhibin-related activities; chemotactic or chemokinetic activities; haemostatic, thrombotic or CC CC thrombolytic activities; receptor or ligand activities; or may be CC involved in oncogenesis, cancer cell proliferation or metastasis. CC Depending on their biological activities, polypeptides and nucleotides of CC the invention are useful for preventing, treating or ameliorating medical conditions, e.g., by protein or gene therapy. Such conditions include CC CC cancers, haematopoietic disorders (e.g., myeloid or lymphoid cell disorders), chronic inflammatory conditions (e.g., asthma or arthritis), CC CC proliferative retinopathy, atherosclerosis, coronary heart disease, arterial ischaemia, bone disorders (e.g., osteoporosis), and abnormal CC CC vascular growth. Polypeptides involved with tissue regeneration and repair (or nucleic acids encoding them) may be used to promote wound CC

healing (e.g., of burns, incisions and ulcers), while those with

CC

```
bacterial and fungal infections in addition to immune disorders.
CC
   Polypeptides with growth factor activity may be used in cell cultures to
CC
   promote cell growth. For example, such polypeptides may be used to
CC
   manipulate stem cells in culture to give rise to neuroepithelial cells
CC
   that can be used to augment or replace cells damaged by illness,
CC
   autoimmune disease or accidental damage. The polypeptides and nucleotides
CC
   may also be used in the diagnosis of the above conditions, and in drug
CC
   screening techniques. The present sequence represents a cDNA encoding a
CC
CC
   novel human polypeptide of the invention.
XX
   Sequence 906 BP; 220 A; 225 C; 216 G; 245 T; 0 other;
SO
                   20.2%; Score 699.2; DB 22;
 Query Match
                                        Length 906;
 Best Local Similarity
                  98.9%; Pred. No. 4.6e-129;
 Matches 704; Conservative
                        0; Mismatches
                                     8;
                                        Indels
                                                        0;
                                                  Gaps
       515 ATGGCTCTGAGTGGAAACTGTAGTCGTTATTATCCTCGAGAACAAGGGTCCGCAGTTCCC 574
Qу
          Db
       906 ATGGCTCTGAGTGGAAACTGTAGTCGTTATTATCCTCGAGAACAAGGGTCCGCAGTTCCC 847
       575 AACTCCTTCCCTGAGGTGGTAGAGCTGAATGTCGGGGGGTCAAGTTTATTTTACTCGCCAT 634
Qу
          846 AACTCCTTCCCTGAGGTGGTAGAGCTGAATGTCGGGGGGTCAAGTTTATTTTACTCGCCAT 787
Db
       635 TCCACATTGATAAGCATCCCTCATTCCCTCCTGTGGAAAATGTTTTCCCCAAAGAGAGAC 694
Qу
          Db
       786 TCCACATTGATAAGCATCCCTCATTCCCTCCTGTGGAAAATGTTTTCCCCAAAGAGAGAC 727
       695 ACGGCTAATGATCTAGCCAAGGACTCCAAGGGAAGGTTTTTCATTGACAGAGATGGATTC 754
Qу
          726 ACGGCTAATGATCTAGCCAAGGACTCCAAGGGAAGGTTTTTCATTGACAGAGATGGATTC 667
Db
       Qу
          Db
       815 CCAGAAAAAGGAAGACTGAAAAGGGAAGCTGAATACTTCCAGCTCCCAGACTTGGTCAAA 874
Qу
          Db
       606 CCAGAAAAAGGAAGACTGAAAAGGGAAGCTGAATACTTCCAGCTCCCAGACTTGGTCAAA 547
       875 CTCCTGACCCCGATGAAATCAAGCAAAGCCCAGATGAATTCTGCCACAGTGACTTTGAA 934
Qу
          Db
       546 CTCCTGACCCCGATGAAATCAAGCAAAGCCCAGATGAATTCTGCCACAGTGACTTTGAA 487
       Qу
          Db
       486 GATGCCTCCCAAGGAAGCGACACAAGAATCTGCCCCCCTTCCTCCCTGCTCCCTGCCGAC 427
       995 CGCAAGTGGGGTTTCATTACTGTGGGTTACAGAGGATCCTGCACCTTGGGCAGAGAGGGA 1054
Qу
          Db
       426 CGCAAGTGGGGTTTCATTACTGTGGGTTACAGAGGATCCTGCACCTTGGGCAGAGAGGGA 367
      1055 CAGGCAGATGCCAAGTTTCGGAGAGTTCCCCGGATTTTGGTTTGTGGAAGGATTTCCTTG 1114
Qу
          Db
       366 CAGGCAGATGCCAAGTTTCGGAGAGTTCCCCGGATTTTGGTTTGTGGAAGGATTTCCTTG 307
```

immunomodulatory activities may be used in the treatment of viral,

CC

```
1115 GCAAAAGAAGTCTTTGGAGAAACTTTGAATGAAAGCAGAGACCCTGATCGAGCCCCAGAA 1174
Qу
             Db
         306 GCAAAAGAAGTCTTTGGAGAAACTTTGAATGAAAGCAGAGCCCTGATCGAGCCCCAGAA 247
        1175 AGATACACCTCCAGATTTTATCTCAAATTCAAGCACCTGGAAAGGGCTTTTG 1226
Qy
             Db
         246 AGATACACCTCCAGATTTTATCTCAAATTCAAGCACCTAATGGGGGCACCTG 195
RESULT 5
AAS34230
TD
    AAS34230 standard; cDNA; 440 BP.
XX
AC
    AAS34230;
XX
DT
    17-DEC-2001 (first entry)
XX
DΕ
    Human cDNA encoding a novel foetal antigen, SEQ ID No 754.
XX
KW
    Human; foetal tissue antigen; ss; antiinflammatory; neuroprotective;
KW
     immunomodulator; cardiovascular; cytostatic; nephrothropic;
KW
     cardiovascular; autoimmune disease; rheumatoid arthritis;
    hyperproliferative disorder; breast neoplasm; cancer;
KW
KW
     cardiovascular disorder; cardiac arrest; cerebrovascular disorder;
ΚW
    cerebral ischaemia; angiogenesis; nervous system disorder;
KW
    Alzheimer's disease; infection; ocular disorder; corneal infection;
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    wound healing; epithelial cell proliferation; food additive.
XX
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    Homo sapiens.
XX
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05-JAN-2001; 2001US-0259678.
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     (HUMA-) HUMAN GENOME SCI INC.
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XX
    Rosen CA, Barash SC, Ruben SM;
PI
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    WPI: 2001-488782/53.
DR
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    P-PSDB; AAU21410.
XX
PT
    New polynucleotides and polypeptides for diagnosing, treating,
    preventing or prognosing e.g. diseases or disorders of the nervous,
PT
PT
    musculoskeletal, excretory, gastrointestinal, reproductive, and
PT
    respiratory systems
XX
PS
    Claim 1; SEQ ID No 754; 642pp; English.
XX
CC
    The invention relates to novel nucleic acids encoding novel human foetal
    antigens. The nucleic acids and proteins are used to prevent, treat (e.g.
CC
    by gene therapy) or ameliorate a medical condition in e.g. humans, mice,
CC
CC
    rabbits, goats, horses, cats, dogs, chickens or sheep. They
CC
    are also used in diagnosing a pathological condition or susceptibility
    to a pathological condition. The antibodies to the antigens can also
CC
CC
    be used in alleviating symptoms associated with the disorders and in
CC
    diagnostic immunoassays e.g. radioimmunoassays or enzyme linked
CC
    immunosorbent assays (ELISA). Disorders which are diagnosed or treated
CC
    include autoimmune diseases e.g. rheumatoid arthritis,
CC
    hyperproliferative disorders e.g. neoplasms of the breast or liver,
CC
    cardiovascular disorders e.g. cardiac arrest, cerebrovascular disorders
CC
    e.q. cerebral ischaemia, angiogenesis, nervous system disorders e.q.
CC
    Alzheimer's disease, infections caused by bacteria, viruses and fungi
CC
    and ocular disorders e.g. corneal infection. The polypeptides can also
    be used to aid wound healing and epithelial cell proliferation, to
CC
CC
    prevent skin aging due to sunburn, to maintain organs before
CC
    transplantation, for supporting cell culture of primary tissues, to
     regenerate tissues and in chemotaxis. The polypeptides can also be used
CC
CC
    as a food additive or preservative to increase or decrease storage
CC
     capabilities, fat content, lipid, protein, carbohydrate, vitamins,
CC
    minerals, cofactors and other nutritional components. Numerous
CC
     examples of diseases and disorders treated by the nucleic acids and
CC
    proteins are given in the specification. The present sequence
                        12.2%;
                                Score 423.4; DB 22;
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                                                     Length 440;
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                                Pred. No. 1.7e-74;
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            Db
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25-MAY-2001; 2001US-293724P.
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    (INCY-) INCYTE GENOMICS INC.
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    Forsythe IJ, Lu Y, Tang YT, Yue H, Raumann BE, Lal PG, Azimzai Y;
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    Lu DAM, Gandhi AR,
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    Swarnakar A, Yao MG, Ding L, He A, Griffin JA, Sanjanwala MM;
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    Gietzen KJ, Lee EA, Xu Y, Au-Young JK, Das D, Lee SY, Chang H;
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    WPI; 2003-092996/08.
    P-PSDB; AAE32081.
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PT
    New human functional transporters and ion channels (TRICH)
PT
    polypeptides, useful for preparing a composition for diagnosing or
PT
    treating a disease associated with decreased expression or
    overexpression of TRICH e.g. cancer -
PT
XX
PS
    Claim 5; Page 200-201; 204pp; English.
XX
CC
    The invention relates to human transporters and ion channels (TRICH)
CC
    polypeptides and nucleic acid molecules encoding such polypeptides.
CC
    TRICH proteins are useful for preparing compositions for diagnosing or
CC
    treating diseases or conditions associated with decreased expression
CC
    or overexpression of functional TRICH e.g. atherosclerosis or cancer.
CC
    The invention is useful in gene therapy. The present sequence is
CC
    human TRICH cDNA.
XX
    Sequence 2398 BP; 644 A; 588 C; 604 G; 562 T; 0 other;
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Db	876	AGCCGCGACCCGACCGGCAGCCGGAGAAGTACACGTCCCGCTTCTACCTCAAGTTCACC	935
Qу	1208	CACCTGGAAAGGGCTTTTGATATGTTGTCAGAGTGTGGATTCCACATGGTGGCCTGTAAC	1267
Db	936	TACTTGGAGCAGGCCTTTGATCGCCTGTCCGAGGCCGGCTTCCACATGGTGGCGTGTAAC	995
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Db	996	TCCTCGGGCACCGCCGCCTTCGTCAACCAGTACCGCGACGACAAGATCTGGAGCAGCTAC	1055
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Db	1056	ACCGAGTACATTTTCTTCCGACCACCTCAGAAAATAGTATCACCTAAACAAGAACATGAA	1115
Qу	1385	TGCTGCAAGAATGGCAAAGGTGACAAAGAAGGGGGAGAGCGGCACGTCTTGCAATGAC	1441
Db	1116	GATAGGAAACATGACAAAGTCACTGATAAAGGAAGTGAAAGTGGGACTTCCTGTAATGAG	1175
Qy	1442	CTCTCCACATCTAGCTGCGACAGCCAGTCTGAGGCCAGCTCTCCCCAGGAGACGGTCATC	1501
Db	1176	CTCTCCACTTCCAGTTGTGACAGCCATTCAGAGGCAAGCACTCCCCAGGACAACCCATCC	1235
Qу	1502	TGTGGTCCCGTGACACGCCAGACCAACATCCAGACTCTGGACCGTCCCATCAAG	1555
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    cytostatic; tumour; gene; ss.
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    Benson DR, Kalos MD, Lodes MJ, Persing DH, Hepler WT, Jiang Y;
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DR
    WPI; 2002-627435/67.
XX
PT
    New isolated polynucleotide and pancreatic tumor polypeptides, useful
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    for diagnosing, preventing and/or treating cancer, particularly
PT
    pancreatic cancer -
XX
PS
    Claim 1; SEQ ID NO 4467; 300pp + Sequence Listing; English.
XX
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CC
    The invention relates to an isolated polynucleotide (I) comprising: (a)
CC
    any of a group of over 4000 nucleotide sequences (ABV94628-ABV99145);
CC
    (b) complements of (a); (c) sequences consisting of at least 20
    contiguous residues of (a); (d) sequences that hybridize to (a), under
CC
    moderately stringent conditions; (e) sequences having at least 75% or 90%
CC
    identity to (a); or (f) degenerate variants of (a). Polypeptides
CC
CC
    (ABP68596-ABP68637) encoded by (I) and oligonucleotide can be used to
CC
    detect cancer in a patient and compositions comprising polypeptides,
CC
    polynucleotides, antibodies, fusion proteins, T cell populations and
CC
    antigen presenting cells expressing the polypeptide are useful in
CC
    treating pancreatic cancer and stimulating an immune response. The
    polynucleotides can be used as probes or primers for nucleic acid
CC
CC
    hybridisation, in the design and preparation of ribozyme molecules for
CC
    inhibiting expression of the tumour polypeptides and proteins in the
CC
    tumour cells, in vaccines and for gene therapy.
CC
    Note: The sequence data for this patent did not form part of the printed
CC
    specification, but was obtained in electronic format directly from WIPO
CC
    at ftp.wipo.int/pub/published pct sequences.
XX
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    cytostatic; tumour; gene; ss.
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os
    Homo sapiens.
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    30-JAN-2002; 2002WO-US02781.
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    30-JAN-2001; 2001US-265305P.
PR
    31-JAN-2001; 2001US-265682P.
PR
    09-FEB-2001; 2001US-267568P.
PR
PR
    21-MAR-2001; 2001US-278651P.
PR
    28-APR-2001; 2001US-287112P.
PR
    16-MAY-2001; 2001US-291631P.
    12-JUL-2001; 2001US-305484P.
PR
    20-AUG-2001; 2001US-313999P.
PR
    27-NOV-2001; 2001US-333626P.
PR
XX
PA
     (CORI-) CORIXA CORP.
XX
ΡI
    Benson DR, Kalos MD, Lodes MJ, Persing DH, Hepler WT, Jiang Y;
XX
DR
    WPI: 2002-627435/67.
XX
PT
    New isolated polynucleotide and pancreatic tumor polypeptides, useful
PT
    for diagnosing, preventing and/or treating cancer, particularly
PT
    pancreatic cancer -
XX
    Claim 1; SEQ ID NO 564; 300pp + Sequence Listing; English.
PS
XX
CC
    The invention relates to an isolated polynucleotide (I) comprising: (a)
CC
    any of a group of over 4000 nucleotide sequences (ABV94628-ABV99145);
CC
     (b) complements of (a); (c) sequences consisting of at least 20
CC
     contiguous residues of (a); (d) sequences that hybridize to (a), under
CC
    moderately stringent conditions; (e) sequences having at least 75% or 90%
CC
    identity to (a); or (f) degenerate variants of (a). Polypeptides
CC
     (ABP68596-ABP68637) encoded by (I) and oligonucleotide can be used to
CC
    detect cancer in a patient and compositions comprising polypeptides,
CC
    polynucleotides, antibodies, fusion proteins, T cell populations and
CC
    antigen presenting cells expressing the polypeptide are useful in
CC
     treating pancreatic cancer and stimulating an immune response. The
CC
    polynucleotides can be used as probes or primers for nucleic acid
    hybridisation, in the design and preparation of ribozyme molecules for
CC
CC
    inhibiting expression of the tumour polypeptides and proteins in the
CC
    tumour cells, in vaccines and for gene therapy.
CC
    Note: The sequence data for this patent did not form part of the printed
     specification, but was obtained in electronic format directly from WIPO
CC
CC
    at ftp.wipo.int/pub/published pct sequences.
XX
     Sequence 614 BP; 177 A; 87 C; 110 G; 236 T; 4 other;
SO
 Query Match
                          5.8%;
                                 Score 201; DB 24; Length 614;
  Best Local Similarity
                         93.3%;
                                 Pred. No. 2.1e-30;
 Matches 210; Conservative
                                0; Mismatches
                                                 15; Indels
                                                                0; Gaps
                                                                            0;
Qу
         3210 ACCAATAGCTCTTTCATAGACTTGTGCTACAAGAAGGTTAAAAGACCAGTTTTATTTTCA 3269
              Db
           1 ACCAATAGCTCTTTCATAGACTTGTGCTACAAGAAGGTTAAAAGACCAGTTTTATTTTCA 60
```

```
3270 GCATTCCTCATGCATTTCAGTGGTAACCAAAAAATAATTTGTCAATTAATAGTTGTGTGC 3329
Qу
           61 GCATTCCTCATGCATTTCAGTGGTAACCAAAAAATAATTTGTCAATTAATAGTTGTGTGC 120
Db
       Qy
           Db
Qу
       181 ATAAAGGCAATTGGATGATATCTGTAGGAGGAAAACAATGACTAA 225
Dh
RESULT 9
ABT09812
TD
   ABT09812 standard; cDNA; 2052 BP.
XX
AC
   ABT09812;
XX
DT
    05-DEC-2002 (first entry)
XX
DE
    Polynucleotide encoding the K+beta M6 protein SEQ ID No 1.
XX
KW
    Cytostatic; cardiant; neuroprotective; immunomodulator; antimigraine;
KW
    sedative; gynaecological;; potassium channel beta subunit; K+betaM6;
    gastrointestinal; reproductive; neural; sleep; low DNA repair capacity;
KW
KW
    hyperpotassium channel activity; cardiovascular; melatonin synthesis;
KW
    mammary cancer tumourigenesis; pineal gland associated disorder;
    pulmonary disorder; immune disorder; NF-kB activity; migraine headache;
KW
KW
    low free-radical buffering capacity; delayed sleep phase syndrome;
    circadian cycle; melatonin secretion; cancer; gene; ss.
KW
XX
os
    Homo sapiens.
XX
PN
    WO200270727-A2.
XX
PD
    12-SEP-2002.
XX
PF
    21-FEB-2002; 2002WO-US05674.
XX
PR
    21-FEB-2001; 2001US-270132P.
PR
    27-MAR-2001; 2001US-278953P.
XX
PA
    (BRIM ) BRISTOL-MYERS SOUIBB CO.
XX
PΙ
    Feder J, Lee L, Chen J, Jackson DG, Ramanathan C, Siemers N;
PΙ
    Chang H;
XX
DR
    WPI; 2002-713455/77.
DR
    P-PSDB; ABJ10886.
XX
PT
    New polynucleotide encoding human potassium channel beta subunit
PT
    polypeptide, useful for diagnosing, preventing, treating or
PТ
    ameliorating e.g. cancer -
XX
PS
    Claim 1; Fig 1; 332pp; English.
XX
```

```
CC
    channel beta subunit (K+betaM6) polypeptide or its variants. The human
    potassium beta subunit polynucleotide or polypeptide is useful for
CC
    diagnosing, preventing, treating or ameliorating a pathological condition
CC
CC
    such as gastrointestinal, reproductive, neural, sleep, cardiovascular or
    pulmonary disorders, a disorder related to hyperpotassium channel
CC
CC
    activity, an immune disorder related to aberrant NF-kB activity, pineal
CC
    gland associated disorders, migraine headaches, disorders associated with
    aberrant melatonin synthesis and/or release or with low DNA repair
CC
CC
    capacities or low free-radical buffering capacity, delayed sleep phase
CC
    syndrome, aberrations in circadian cycle, mammary cancer tumourigenesis,
CC
    age related disorders associated with decreased melatonin secretion, or
CC
    cancer. This polynucleotide sequence represents the cDNA encoding the
CC
    potassium channel beta subunit (K+betaM6) protein of the invention.
XX
    Sequence 2052 BP; 380 A; 640 C; 607 G; 425 T; 0 other;
SO
 Query Match
                        4.8%;
                              Score 167; DB 24; Length 2052;
 Best Local Similarity
                       64.6%;
                              Pred. No. 1.5e-23;
 Matches 267; Conservative
                             0; Mismatches 140; Indels
                                                          6;
                                                             Gaps
                                                                     1;
         967 CCCCCTTCCTCCCTGCTCCCTGCCGACCGCAAGTGGGGTTTCATTACTGTGGGTTACAG 1026
Qу
                          111
                               705 CACGCCGTCCCAGTCGCTGGACGGCAGCCGGCGCTCGGGCTACATCACCATCGGCTACCG 764
Db
Qу
        1027 AGGATCCTGCACCTTGGGCAGAGAGGGACAGGCAGATGCCAAGTTTCGGAGAGTTCCCCG 1086
             Db
        765 CGGCTCCTACACCATCGGGCGGGACGCGCAGGCGGACGCCAAGTTCCGGCGAGTGGCGCG 824
Qу
        1087 GATTTTGGTTTGTGGAAGGATTTCCTTGGCAAAAGAAGTCTTTGGAGAAACTTTGAATGA 1146
                   Db
         825 CATCACCGTTTGCGGAAAGACGTCGCTGGCCAAGGAGGTGTTTGGGGACACCCTGAACGA 884
        1147 AAGCAGAGCCCTGATCGAGCCCCAGAAAGATACACCTCCAGATTTTATCTCAAATTCAA 1206
Qу
                             1111 | 1111 | 11 | 11
         885 AAGCCGGGACCCCGACCGTCCCCCGGAGCGCTACACCTCGCGCTATTACCTCAAGTTCAA 944
Db
        1207 GCACCTGGAAAGGGCTTTTGATATGTTGTCAGAGTGTGGATTCCACATGGTGGCCTGTAA 1266
Qу
                       945 CTTCCTGGAGCAGGCCTTCGACAAGCTGTCCGAGTCGGGCTTCCACATGGTGGCGTGCAG 1004
Db
        1267 CTCATCGGTGACAGCATCTTT-----CATCAACCAATATACAGATGACAAGATCTGGTC 1320
Qу
            111 111 11
                           1 11
                                    1005 CTCCACGGGCACCTGCGCCTTTGCCAGCAGCCAGCCAGAGCGAGGACAAGATCTGGAC 1064
Db
        1321 AAGCTACACTGAATATGTCTTCTACCGTGAGCCTTCCAGATGGTCACCCTCAC 1373
Qу
             1 | | | | | | |
        1065 CAGCTACACCGAGTACGTCTTCTGCAGGGAGTGAGCTCCCCAGACCCCCTCGC 1117
Db
RESULT 10
ABQ88125/c
ID
    ABQ88125 standard; cDNA; 109201 BP.
XX
AC
    ABQ88125;
XX
DT
    18-SEP-2002 (first entry)
```

The invention relates to an isolated polynucleotide encoding a potassium

CC

```
XX
     Human osteoblast differentiation related cDNA SEQ ID NO 32.
DE
XX
     Human; osteoblast; stem cell differentiation; bone tissue deposition;
KW
KW
     osteoporosis; osteopathic; ss.
XX
os
    Homo sapiens.
XX
PN
    WO200250301-A2.
XX
    27-JUN-2002.
PD
XX
PF
    18-DEC-2001; 2001WO-US48276.
XX
PR
     18-DEC-2000; 2000US-255882P.
PR
     24-APR-2001; 2001US-285691P.
XX
PA
     (GENE-) GENE LOGIC INC.
     (PROC ) PROCTER & GAMBLE CO.
PA
XX
PΙ
     Ji D, Axelrod DW, Cook JS, Jaiswal N, Einstein R, Houghton A;
ΡI
    Mertz L;
XX
DR
    WPI; 2002-557663/59.
XX
PT
    Use of genes and their expression profiles associated with osteoblast
PT
     differentiation for screening modulators bone formation, for diagnosing
PT
     or treating e.g. osteoporosis, or as markers for the differentiation
PT
    process
XX
PS
    Claim 1; SEQ ID NO 32; 78pp + Sequence Listing; English.
XX
CC
    The invention relates to genes and their expression profiles are used
CC
CC
     (a) screening modulators of precursor stem cell differentiation into
CC
     osteoblasts, or bone tissue deposition;
CC
     (b) diagnosing abnormal deposition of bone tissue, abnormal rate of
CC
     osteoblast formation or osteoporosis; or
CC
     (c) treating or monitoring treatment of the conditions cited in (b), or
CC
     monitoring the progression of bone tissue deposition.
CC
     Specific conditions include postmenopausal osteoporosis, glucocorticoid
CC
     osteoporosis or male osteoporosis, osteopenia, osteodystrophy,
CC
     drug-induced abnormalities in bone formation or bone loss, conditions
CC
     that involve altered bone metabolism (e.g. idiopathic juvenile
CC
     osteoporosis), skeletal disease linked to breast cancer, mastocytosis,
CC
     Fanconi syndrome or fibrous dysplasia. The present sequence is that of an
CC
     osteoblast differentiation associated cDNA marker of the invention.
CC
     Note: The sequence data for this patent did not form part of the printed
CC
     specification, but was obtained in electronic format directly from WIPO
CC
     at ftp.wipo.int/pub/published pct sequences.
XX
SO
     Sequence 109201 BP; 32871 A; 23488 C; 22108 G; 30734 T; 0 other;
  Query Match
                           4.8%; Score 167; DB 24; Length 109201;
  Best Local Similarity
                          64.6%;
                                  Pred. No. 3.5e-23;
  Matches 267; Conservative
                                 0; Mismatches 140; Indels
                                                                 6; Gaps
                                                                              1;
```

```
967 CCCCCCTTCCTCCTGCTGCCGACCGCAAGTGGGGTTTCATTACTGTGGGTTACAG 1026
Qy
                       9291 CACGCCGTCCCAGTCGCTGGACGGCAGCCGGCGCTCGGGCTACATCACCATCGGCTACCG 9232
Db
       1027 AGGATCCTGCACCTTGGGCAGAGAGGGACAGGCAGATGCCAAGTTTCGGAGAGTTCCCCG 1086
Qу
            9231 CGGCTCCTACACCATCGGGCGGGACGCGCAGGCGGACGCCAAGTTCCGGCGAGTGGCGCG 9172
Db
Qy
       1087 GATTTTGGTTTGTGGAAGGATTTCCTTGGCAAAAGAAGTCTTTGGAGAAACTTTGAATGA 1146
                 9171 CATCACCGTTTGCGGAAAGACGTCGCTGGCCAAGGAGGTGTTTGGGGACACCCTGAACGA 9112
Db
Qy
       1147 AAGCAGAGCCCTGATCGAGCCCCAGAAAGATACACCTCCAGATTTTATCTCAAATTCAA 1206
           9111 AAGCCGGGACCCGACCGTCCCCGGAGCGCTACACCTCGCGCTATTACCTCAAGTTCAA 9052
Db
Qу
       1207 GCACCTGGAAAGGGCTTTTGATATGTTGTCAGAGTGTGGATTCCACATGGTGGCCTGTAA 1266
                     Db
       9051 CTTCCTGGAGCAGGCCTTCGACAAGCTGTCCGAGTCGGGCTTCCACATGGTGGCGTGCAG 8992
Qу
       1267 CTCATCGGTGACAGCATCTTT-----CATCAACCAATATACAGATGACAAGATCTGGTC 1320
           1 11
                                 11 11 1 1 1 1 1 11 11 11 11 11 1
Db
       1321 AAGCTACACTGAATATGTCTTCTACCGTGAGCCTTCCAGATGGTCACCCTCAC 1373
Qу
            - 1
                                             Dh
       8931 CAGCTACACCGAGTACGTCTTCTGCAGGGAGTGAGCTCCCCAGACCCCCTCGC 8879
RESULT 11
ABQ40654
   ABQ40654 standard; DNA; 854 BP.
XX
AC
   ABO40654;
XX
\mathbf{DT}
    12-JUL-2002 (first entry)
XX
DΕ
    Oligonucleotide for detecting cytosine methylation SEQ ID NO 27245.
XX
KW
    Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
    drug; side effect; cancer; central nervous system; cardiovascular;
KW
KW
    gastrointestinal; respiratory system; single nucleotide polymorphism;
    SNP; cell differentiation; ds.
KW
XX
OS
    Homo sapiens.
XX
PN
    WO200218632-A2.
XX
PD
    07-MAR-2002.
XX
PF
    01-SEP-2001; 2001WO-EP10074.
XX
PR
    01-SEP-2000; 2000DE-1043826.
    05-SEP-2000; 2000DE-1044543.
PR
XX
PA
    (EPIG-) EPIGENOMICS AG.
XX
```

```
Olek A, Piepenbrock C, Berlin K, Guetig D;
PΙ
XX
    WPI; 2002-371829/40.
DR
XX
PT
    Determining the degree of cytosine methylation in genomic DNA, useful
     for diagnosis and prognosis, comprises selective hybridization of
PT
    amplicons from chemically treated DNA -
PT
XX
PS
    Claim 12; 56pp + Sequence Listing; 56pp; German.
XX
CC
    This invention describes a novel method for determining the degree of
    methylation of a particular cytosine in a motif 5'-CpG-3', present in a
CC
CC
    genomic sample of DNA. The sample is treated chemically to convert
CC
    cytosine (C) but not methylated C, to uracil, then part of the genomic
CC
    DNA that contains the target C is amplified to form a labeled amplicon.
CC
    The amplicon is hybridised to two classes, each with at least one
CC
    member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers
CC
    and the degree of hybridisation to both classes is determined from the
    label on the amplicon. From the ratio of labels hybridised to the two
CC
CC
    classes of oligomers, the degree of methylation is calculated. The method
CC
    is used: (i) for diagnosis and/or prognosis of side effects of
CC
    therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders
CC
    of the central nervous, cardiovascular, gastrointestinal and respiratory
     systems etc., particularly by detecting mutations or single nucleotide
CC
CC
    polymorphisms (SNP's); and (ii) for differentiation of cell or tissue
CC
    types and for investigating cell differentiation. The method allows the
CC
    methylation status of many C residues to be determined simultaneously.
CC
    ABQ13410-ABQ54121 represent genomic DNA sequences used to illustrate the
CC
    method for determining the degree of cytosine methylation described in
CC
    the disclosure of the invention.
XX
     Sequence 854 BP; 131 A; 98 C; 289 G; 336 T; 0 other;
SQ
 Query Match
                         3.3%; Score 114.2; DB 24; Length 854;
  Best Local Similarity
                        59.7%;
                                Pred. No. 3.6e-13;
 Matches 213; Conservative
                               0; Mismatches 138; Indels
                                                             6;
                                                                Gaps
                                                                        1;
Qу
        1001 TGGGGTTTCATTACTGTGGGTTACAGAGGATCCTGCACCTTGGGCAGAGAGGGACAGGCA 1060
             Db
         491 TCGGGTTATATTATTATCGGTTATCGCGGTTTTTATATTATCGGGCGGACGCGTAGGCG 550
        1061 GATGCCAAGTTTCGGAGAGTTCCCCGGATTTTGGTTTGTGGAAGGATTTCCTTGGCAAAA 1120
Qу
             551 GACGTTAAGTTTCGGCGAGTGGCGCGTATTATCGTTTGCGGAAAGACGTCGTTGGTTAAG 610
Db
        1121 GAAGTCTTTGGAGAAACTTTGAATGAAAGCAGAGCCCTGATCGAGCCCCAGAAAGATAC 1180
Qу
             1 11 1 11
         611 GAGGTGTTTGGGGGATATTTTGAACGAAAGTCGGGATTTCGATCGTTTTTCGGAGCGTTAT 670
Db
Qу
        1181 ACCTCCAGATTTTATCTCAAATTCAAGCACCTGGAAAGGGCTTTTGATATGTTGTCAGAG 1240
               ++++
                                                671 ATTTCGCGTTATTATTTTAAGTTTAATTTTTTGGAGTAGGTTTTCGATAAGTTGTTCGAG 730
Db
Qу
        1241 TGTGGATTCCACATGGTGGCCTGTAACTCATCGGTGACAGCATCTTT-----CATCAAC 1294
```

731 TCGGGTTTTTATATGGTGGCGTGTAGTTTTACGGGTATTTGCGTTTTTGTTAGTAGTATC 790

Db

 $\Pi\Pi$

```
RESULT 12
ABQ40655/c
     ABQ40655 standard; DNA; 854 BP.
ID
XX
AC
     ABO40655;
XX
DT
     12-JUL-2002 (first entry)
XX
DΕ
     Oligonucleotide for detecting cytosine methylation SEQ ID NO 27246.
XX
     Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
KW
KW
     drug; side effect; cancer; central nervous system; cardiovascular;
KW
     gastrointestinal; respiratory system; single nucleotide polymorphism;
KW
     SNP; cell differentiation; ds.
XX
OS
     Homo sapiens.
XX
PN
     WO200218632-A2.
XX
PD
     07-MAR-2002.
XX
PF
     01-SEP-2001; 2001WO-EP10074.
XX
PR
     01-SEP-2000; 2000DE-1043826.
     05-SEP-2000; 2000DE-1044543.
PR
XX
     (EPIG-) EPIGENOMICS AG.
PA
XX
PΙ
     Olek A, Piepenbrock C, Berlin K, Guetig D;
XX
     WPI; 2002-371829/40.
DR
XX
PT
     Determining the degree of cytosine methylation in genomic DNA, useful
PT
     for diagnosis and prognosis, comprises selective hybridization of
PT
     amplicons from chemically treated DNA -
XX
PS
     Claim 12; 56pp + Sequence Listing; 56pp; German.
XX
CC
     This invention describes a novel method for determining the degree of
CC
```

This invention describes a novel method for determining the degree of methylation of a particular cytosine in a motif 5'-CpG-3', present in a genomic sample of DNA. The sample is treated chemically to convert cytosine (C) but not methylated C, to uracil, then part of the genomic DNA that contains the target C is amplified to form a labeled amplicon. The amplicon is hybridised to two classes, each with at least one member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers and the degree of hybridisation to both classes is determined from the label on the amplicon. From the ratio of labels hybridised to the two classes of oligomers, the degree of methylation is calculated. The method is used: (i) for diagnosis and/or prognosis of side effects of therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders of the central nervous, cardiovascular, gastrointestinal and respiratory systems etc., particularly by detecting mutations or single nucleotide

CC

```
polymorphisms (SNP's); and (ii) for differentiation of cell or tissue
CC
    types and for investigating cell differentiation. The method allows the
CC
    methylation status of many C residues to be determined simultaneously.
    ABQ13410-ABQ54121 represent genomic DNA sequences used to illustrate the
CC
    method for determining the degree of cytosine methylation described in
    the disclosure of the invention.
CC
XX
    Sequence 854 BP; 336 A; 289 C; 98 G; 131 T; 0 other;
SO
 Query Match
                       3.3%; Score 114.2; DB 24; Length 854;
 Best Local Similarity
                      59.7%; Pred. No. 3.6e-13;
 Matches 213; Conservative
                            0; Mismatches 138; Indels
                                                        6; Gaps
       1001 TGGGGTTTCATTACTGTGGGTTACAGAGGATCCTGCACCTTGGGCAGAGAGGGACAGGCA 1060
Qy
            Db
        1061 GATGCCAAGTTTCGGAGAGTTCCCCGGATTTTGGTTTGTGGAAGGATTTCCTTGGCAAAA 1120
Qу
            304 GACGTTAAGTTTCGGCGAGTGGCGCGTATTATCGTTTGCGGAAAGACGTCGTTGGTTAAG 245
Db
       1121 GAAGTCTTTGGAGAAACTTTGAATGAAAGCAGAGACCCTGATCGAGCCCCAGAAAGATAC 1180
Qv
            11111
                                                      244 GAGGTGTTTGGGGATATTTTGAACGAAAGTCGGGATTTCGATCGTTTTTCGGAGCGTTAT 185
Db
       1181 ACCTCCAGATTTTATCTCAAATTCAAGCACCTGGAAAGGGCTTTTGATATGTTGTCAGAG 1240
Qу
            +1111
                                            184 ATTTCGCGTTATTATTTTAAGTTTAATTTTTTGGAGTAGGTTTTCGATAAGTTGTTCGAG 125
Db
       1241 TGTGGATTCCACATGGTGGCCTGTAACTCATCGGTGACAGCATCTTT-----CATCAAC 1294
Qу
              111
Db
        124 TCGGGTTTTTATATGGTGGCGTGTAGTTTTACGGGTATTTGCGTTTTTGTTAGTAGTATC 65
       1295 CAATATACAGATGACAAGATCTGGTCAAGCTACACTGAATATGTCTTCTACCGTGAG 1351
Qу
             1 11 1 14 11 11111 111
                                   64 GATTAGAGCGAGGATAAGATTTGGATTAGTTATATCGAGTACGTTTTTTTGTAGGGAG 8
Db
RESULT 13
ABQ13668
    ABQ13668 standard; DNA; 1757 BP.
XX
AC
    ABQ13668;
XX
DT
    12-JUL-2002 (first entry)
XX
DE
    Oligonucleotide for detecting cytosine methylation SEQ ID NO 259.
XX
KW
    Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
KW
    drug; side effect; cancer; central nervous system; cardiovascular;
KW
    gastrointestinal; respiratory system; single nucleotide polymorphism;
KW
    SNP; cell differentiation; ds.
XX
OS
    Homo sapiens.
XX
PN
    W0200218632-A2.
XX
```

```
07-MAR-2002.
PD
XX
    01-SEP-2001; 2001WO-EP10074.
PF
XX
PR
    01-SEP-2000; 2000DE-1043826.
    05-SEP-2000; 2000DE-1044543.
PR
XX
PA
    (EPIG-) EPIGENOMICS AG.
XX
PΙ
    Olek A, Piepenbrock C,
                           Berlin K, Guetig D;
XX
DR
    WPI; 2002-371829/40.
XX
    Determining the degree of cytosine methylation in genomic DNA, useful
PT
PT
    for diagnosis and prognosis, comprises selective hybridization of
PT
    amplicons from chemically treated DNA
XX
PS
    Claim 12; 56pp + Sequence Listing; 56pp; German.
XX
CC
    This invention describes a novel method for determining the degree of
CC
    methylation of a particular cytosine in a motif 5'-CpG-3', present in a
    genomic sample of DNA. The sample is treated chemically to convert
CC
CC
    cytosine (C) but not methylated C, to uracil, then part of the genomic
CC
    DNA that contains the target C is amplified to form a labeled amplicon.
    The amplicon is hybridised to two classes, each with at least one
CC
CC
    member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers
CC
    and the degree of hybridisation to both classes is determined from the
CC
    label on the amplicon. From the ratio of labels hybridised to the two
CC
    classes of oligomers, the degree of methylation is calculated. The method
CC
    is used: (i) for diagnosis and/or prognosis of side effects of
CC
    therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders
CC
    of the central nervous, cardiovascular, gastrointestinal and respiratory
CC
    systems etc., particularly by detecting mutations or single nucleotide
    polymorphisms (SNP's); and (ii) for differentiation of cell or tissue
CC
CC
    types and for investigating cell differentiation. The method allows the
CC
    methylation status of many C residues to be determined simultaneously.
CC
    ABQ13410-ABQ54121 represent genomic DNA sequences used to illustrate the
CC
    method for determining the degree of cytosine methylation described in
CC
    the disclosure of the invention.
XX
    Sequence 1757 BP; 246 A; 209 C; 640 G; 662 T; 0 other;
SO
 Query Match
                         3.2%;
                               Score 109.8; DB 24;
                                                    Length 1757;
 Best Local Similarity
                        62.6%;
                               Pred. No. 3.1e-12;
 Matches 171; Conservative
                              0; Mismatches 102;
                                                   Indels
                                                             0; Gaps
                                                                        0;
        1001 TGGGGTTTCATTACTGTGGGTTACAGAGGATCCTGCACCTTGGGCAGAGAGGGACAGGCA 1060
Qу
             1 11
                                                       1 11 1
        Db
Qу
        1061 GATGCCAAGTTTCGGAGAGTTCCCCGGATTTTGGTTTGTGGAAGGATTTCCTTGGCAAAA 1120
             Db
        1545 GACGTTAAGTTTCGGCGAGTGGCGCGTATTATCGTTTGCGGAAAGACGTCGTTGGTTAAG 1604
Qу
        1121 GAAGTCTTTGGAGAAACTTTGAATGAAAGCAGAGACCCTGATCGAGCCCCAGAAAGATAC 1180
             \Box
                                                           1 11 1 11
Db
        1605 GAGGTGTTTGGGGGATATTTTGAACGAAAGTCGGGATTTCGATCGTTTTTCGGAGCGTTAT 1664
```

```
1181 ACCTCCAGATTTTATCTCAAATTCAAGCACCTGGAAAGGGCTTTTGATATGTTGTCAGAG 1240
Qу
               1665 ATTTCGCGTTATTATTTTAAGTTTAATTTTTTGGAGTAGGTTTTCGATAAGTTGTTCGAG 1724
Db
        1241 TGTGGATTCCACATGGTGGCCTGTAACTCATCG 1273
Qγ
             Db
        1725 TCGGGTTTTTATATGGTGGCGTGTAGTTTTACG 1757
RESULT 14
ABQ13669/c
    ABQ13669 standard; DNA; 1757 BP.
XX
AC
    ABQ13669;
XX
DT
    12-JUL-2002 (first entry)
XX
DE
    Oligonucleotide for detecting cytosine methylation SEQ ID NO 260.
XX
KW
    Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
KW
    drug; side effect; cancer; central nervous system; cardiovascular;
    gastrointestinal; respiratory system; single nucleotide polymorphism;
KW
KW
    SNP; cell differentiation; ds.
XX
OS
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PR
    05-SEP-2000; 2000DE-1044543.
XX
PΑ
     (EPIG-) EPIGENOMICS AG.
XX
    Olek A, Piepenbrock C, Berlin K, Guetig D;
PΙ
XX
DR
    WPI; 2002-371829/40.
XX
PT
    Determining the degree of cytosine methylation in genomic DNA, useful
PT
    for diagnosis and prognosis, comprises selective hybridization of
PΤ
    amplicons from chemically treated DNA
XX
PS
    Claim 12; 56pp + Sequence Listing; 56pp; German.
XX
CC
    This invention describes a novel method for determining the degree of
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    methylation of a particular cytosine in a motif 5'-CpG-3', present in a
CC
    genomic sample of DNA. The sample is treated chemically to convert
CC
    cytosine (C) but not methylated C, to uracil, then part of the genomic
CC
    DNA that contains the target C is amplified to form a labeled amplicon.
CC
    The amplicon is hybridised to two classes, each with at least one
CC
    member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers
CC
    and the degree of hybridisation to both classes is determined from the
CC
    label on the amplicon. From the ratio of labels hybridised to the two
```

```
classes of oligomers, the degree of methylation is calculated. The method
CC
    is used: (i) for diagnosis and/or prognosis of side effects of
CC
    therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders
CC
    of the central nervous, cardiovascular, gastrointestinal and respiratory
CC
    systems etc., particularly by detecting mutations or single nucleotide
CC
    polymorphisms (SNP's); and (ii) for differentiation of cell or tissue
CC
CC
    types and for investigating cell differentiation. The method allows the
    methylation status of many C residues to be determined simultaneously.
CC
CC
    ABQ13410-ABQ54121 represent genomic DNA sequences used to illustrate the
CC
    method for determining the degree of cytosine methylation described in
CC
    the disclosure of the invention.
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DT
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    Oligonucleotide for detecting cytosine methylation SEQ ID NO 27247.
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KW
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KW
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KW
    SNP; cell differentiation; ds.
XX
OS
    Homo sapiens.
XX
PN
    WO200218632-A2.
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XX
     07-MAR-2002.
PD
XX
    01-SEP-2001; 2001WO-EP10074.
PF
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PR
    01-SEP-2000; 2000DE-1043826.
    05-SEP-2000; 2000DE-1044543.
PR
XX
PA
     (EPIG-) EPIGENOMICS AG.
XX
PΙ
    Olek A, Piepenbrock C,
                            Berlin K,
                                       Guetiq D;
XX
DR
    WPI: 2002-371829/40.
XX
PT
    Determining the degree of cytosine methylation in genomic DNA, useful
     for diagnosis and prognosis, comprises selective hybridization of
PT
PT
     amplicons from chemically treated DNA -
XX
PS
    Claim 12; 56pp + Sequence Listing; 56pp; German.
XX
CC
    This invention describes a novel method for determining the degree of
    methylation of a particular cytosine in a motif 5'-CpG-3', present in a
CC
CC
    genomic sample of DNA. The sample is treated chemically to convert
CC
     cytosine (C) but not methylated C, to uracil, then part of the genomic
    DNA that contains the target C is amplified to form a labeled amplicon.
CC
    The amplicon is hybridised to two classes, each with at least one
CC
CC
    member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers
CC
    and the degree of hybridisation to both classes is determined from the
CC
    label on the amplicon. From the ratio of labels hybridised to the two
CC
     classes of oligomers, the degree of methylation is calculated. The method
CC
    is used: (i) for diagnosis and/or prognosis of side effects of
    therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders
CC
CC
    of the central nervous, cardiovascular, gastrointestinal and respiratory
CC
     systems etc., particularly by detecting mutations or single nucleotide
CC
    polymorphisms (SNP's); and (ii) for differentiation of cell or tissue
CC
    types and for investigating cell differentiation. The method allows the
CC
    methylation status of many C residues to be determined simultaneously.
CC
    ABQ13410-ABQ54121 represent genomic DNA sequences used to illustrate the
CC
    method for determining the degree of cytosine methylation described in
     the disclosure of the invention.
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Search completed: January 28, 2004, 21:07:41 Job time: 872 secs

GenCore version 5.1.6 Copyright (c) 1993 - 2004 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 28, 2004, 20:50:10; Search time 192 Seconds

(without alignments)

7972.483 Million cell updates/sec

Title: US-10-056-884A-1

Perfect score: 3468

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 569978 seqs, 220691566 residues

Total number of hits satisfying chosen parameters: 1139956

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database: Issued Patents NA:*

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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	2	73.8	2.1	1091	4	US-09-328-965-1	Sequence 1, Appli
С	3	73.4	2.1	4055	4	US-09-620-312D-706	Sequence 706, App
	4	71.2	2.1	1701	4	US-09-996-243-114	Sequence 114, App
	5	69.2	2.0	1441	3	US-08-821-994-63	Sequence 63, Appl
	6	69.2	2.0	2246	4	US-09-363-708-3	Sequence 3, Appli
	7	69.2	2.0	2246	4	US-09-083-587-3	Sequence 3, Appli
	8	69.2	2.0	2406	4	US-09-594-506-37	Sequence 37, Appl
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	29	67.2	1.9	2320	4	US-09-202-904A-13	Sequence 13, Appl
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	34	66.8	1.9	3238	5	PCT-US94-10080-5	Sequence 5, Appli
	35	66.6	1.9	1507	4	US-09-453-323-1	Sequence 1, Appli
	36	66.6	1.9	3334	4	US-09-996-243-288	Sequence 288, App
	37	66.6	1.9	5173	1	US-08-242-677-1	Sequence 1, Appli
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	40	66.4	1.9	2665	4	US-08-971-089-5	Sequence 5, Appli
	41	66.4	1.9	2718	4	US-09-667-135-1	Sequence 1, Appli
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	44	66.2	1.9	2136	4	US-09-996-243-302	Sequence 302, App
	45	66.2	1.9	2218	4	US-09-016-434-1157	Sequence 1157, Ap

ALIGNMENTS

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; APPLICANT: Rosen et al.
  TITLE OF INVENTION: 49 Human Secreted Proteins
  FILE REFERENCE: PZ032P1
  CURRENT APPLICATION NUMBER: US/09/904,615
  CURRENT FILING DATE: 2001-07-16
  PRIOR APPLICATION NUMBER: 09/511,554
  PRIOR FILING DATE: 2000-02-23
  PRIOR APPLICATION NUMBER: 60/097,917
; PRIOR FILING DATE: 1998-08-25
  PRIOR APPLICATION NUMBER: 60/098,634
  PRIOR FILING DATE: 1998-08-31
; NUMBER OF SEQ ID NOS: 170
  SOFTWARE: PatentIn Ver. 2.0
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   LOCATION: (63)
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; Patent No. 6501008
; GENERAL INFORMATION:
; APPLICANT: Nevins, Donald J.
; APPLICANT: Simmons, Carl
 APPLICANT: The Regents of the University of California
  TITLE OF INVENTION: Endo- and Exo-Glucanases and Gene
  FILE REFERENCE: 023070-096600US
  CURRENT APPLICATION NUMBER: US/09/328,965
  CURRENT FILING DATE: 1999-06-09
  EARLIER APPLICATION NUMBER: US 60/088,780
 EARLIER FILING DATE: 1998-06-10
  NUMBER OF SEQ ID NOS: 3
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   NAME/KEY: CDS
   LOCATION: (68)..(979)
   OTHER INFORMATION: endo-1,3;1,4-beta glucanase
US-09-328-965-1
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          Db
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                Db
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          Db
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US-09-620-312D-706/c
; Sequence 706, Application US/09620312D
; Patent No. 6569662
; GENERAL INFORMATION:
 APPLICANT: Tang, Y. Tom
APPLICANT: Liu, Chenghua
 APPLICANT: Asundi, Vinod
 APPLICANT: Zhang, Jie
 APPLICANT: Ren, Feiyan
 APPLICANT: Chen, Rui-hong
 APPLICANT: Zhao, Qing A.
 APPLICANT: Wehrman, Tom
 APPLICANT: Xue, Aidong J.
; APPLICANT: Yang, Yonghong
; APPLICANT: Wang, Jian-Rui
; APPLICANT: Zhou, Ping
; APPLICANT: Ma, Yunging
; APPLICANT: Wang, Dunrui
; APPLICANT: Wang, Zhiwei
 APPLICANT: John Tillinghast APPLICANT: Drmanac, Radoje T.
 TITLE OF INVENTION: No. 6569662el Nucleic Acids and
 TITLE OF INVENTION: Polypeptides
; FILE REFERENCE: 784CIP2B
 CURRENT APPLICATION NUMBER: US/09/620,312D
 CURRENT FILING DATE: 2000-07-19
 PRIOR APPLICATION NUMBER: 09/552,317
 PRIOR FILING DATE: 2000-04-25
 PRIOR APPLICATION NUMBER: 09/488,725
 PRIOR FILING DATE: 2000-01-21
  NUMBER OF SEQ ID NOS: 1105
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  TYPE: DNA
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  FEATURE:
  NAME/KEY: CDS
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; Sequence 114, Application US/09996243
; Patent No. 6478825
; GENERAL INFORMATION:
 APPLICANT: Ashkenazi, Avi J.
  APPLICANT: Baker, Kevin P.
  APPLICANT:
          Botstein, David
  APPLICANT: Desnoyers, Luc
  APPLICANT: Eaton, Dan L.
  APPLICANT: Ferrara, Napoleone
  APPLICANT: Fong, Sherman
  APPLICANT: Gerber, Hanspeter
  APPLICANT:
           Gerritsen, Mary E.
  APPLICANT:
           Goddard, Audrey
  APPLICANT:
           Godowski, Paul J.
  APPLICANT: Grimaldi, J. Christopher
  APPLICANT: Gurney, Austin L.
  APPLICANT: Kljavin, Ivar J.
 APPLICANT: Napier, Mary A.
 APPLICANT: Pan, James
  APPLICANT: Paoni, Nicholas F.
  APPLICANT: Roy, Margaret Ann
  APPLICANT: Stewart, Timothy A.
  APPLICANT: Tumas, Daniel
  APPLICANT: Watanabe, Colin K.
  APPLICANT:
           Williams, P. Mickey
  APPLICANT:
           Wood, William I.
  APPLICANT:
           Zhang, Zemin
  TITLE OF INVENTION: Secreted and Transmembrane Polypeptides and Nucleic
  TITLE OF INVENTION: Acids Encoding the Same
  FILE REFERENCE: P2730P1C13
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CURRENT APPLICATION NUMBER: US/09/996,243

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; CURRENT FILING DATE: 2001-11-14
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- ; PRIOR APPLICATION NUMBER: 60/049787
- ; PRIOR FILING DATE: 1997-06-16
- ; PRIOR APPLICATION NUMBER: 60/062250
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- PRIOR FILING DATE: 1997-11-12
- ; PRIOR APPLICATION NUMBER: 60/065311
- ; PRIOR FILING DATE: 1997-11-13
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- ; PRIOR FILING DATE: 1997-11-24
- ; PRIOR APPLICATION NUMBER: 60/075945
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- ; PRIOR APPLICATION NUMBER: 60/087106
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- ; PRIOR APPLICATION NUMBER: 60/087607
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- ; PRIOR FILING DATE: 1998-06-05
- ; PRIOR APPLICATION NUMBER: 60/088217
- ; PRIOR FILING DATE: 1998-06-05
- ; PRIOR APPLICATION NUMBER: 60/088655
- ; PRIOR FILING DATE: 1998-06-09
- ; PRIOR APPLICATION NUMBER: 60/088734
- ; PRIOR FILING DATE: 1998-06-10

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- ; PRIOR APPLICATION NUMBER: 60/088824
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- ; PRIOR FILING DATE: 1998-06-18
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- ; PRIOR APPLICATION NUMBER: 60/089947
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 PRIOR FILING DATE: 1998-07-02
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PRIOR FILING DATE: 1998-07-07
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PRIOR APPLICATION NUMBER: 60/092182

PRIOR FILING DATE: 1998-07-09

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US-08-821-994-63
; Sequence 63, Application US/08821994A
; Patent No. 6228643
; GENERAL INFORMATION:
; APPLICANT: Greenland, Andrew J
 APPLICANT: Thomas, Didier RP
 APPLICANT: Jepson, Ian
 TITLE OF INVENTION: Promoters
; FILE REFERENCE: PPD 50108
; CURRENT APPLICATION NUMBER: US/08/821,994A
; CURRENT FILING DATE: 1997-03-22
; EARLIER APPLICATION NUMBER: PCT/GB97/00729
; EARLIER FILING DATE: 1997-03-18
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; Patent No. 6399747
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   APPLICANT: Schmandt, et al.
   TITLE OF INVENTION: NOVEL SHC BINDING PROTEIN
   NUMBER OF SEQUENCES: 12
   CORRESPONDENCE ADDRESS:
     ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun
     STREET: 233 South Wacker Drive/6300 Sears Tower
   CITY: Chicago
     STATE: Illinois
     COUNTRY: United States of America
     ZIP: 60606-6402
   COMPUTER READABLE FORM:
     MEDIUM TYPE: Floppy disk
     COMPUTER: IBM PC compatible
     OPERATING SYSTEM: PC-DOS/MS-DOS
     SOFTWARE: PatentIn Release #1.0, Version #1.30
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     APPLICATION NUMBER: US/09/363,708
     FILING DATE:
     CLASSIFICATION:
   ATTORNEY/AGENT INFORMATION:
     NAME: Clough, David W.
     REGISTRATION NUMBER: 36,107
     REFERENCE/DOCKET NUMBER: 01017/34451
    TELECOMMUNICATION INFORMATION:
     TELEPHONE: (312) 474-6300
     TELEFAX: (312) 474-0448
  INFORMATION FOR SEQ ID NO: 3:
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     TOPOLOGY: linear
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   APPLICANT: Schmandt, et al.
   TITLE OF INVENTION: NOVEL SHC BINDING PROTEIN
   NUMBER OF SEQUENCES: 12
   CORRESPONDENCE ADDRESS:
;
     ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun
     STREET: 233 South Wacker Drive/6300 Sears Tower
     CITY: Chicago
     STATE: Illinois
     COUNTRY: United States of America
     ZIP: 60606-6402
   COMPUTER READABLE FORM:
     MEDIUM TYPE: Floppy disk
     COMPUTER: IBM PC compatible
     OPERATING SYSTEM: PC-DOS/MS-DOS
     SOFTWARE: PatentIn Release #1.0, Version #1.30
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     FILING DATE:
     CLASSIFICATION:
   ATTORNEY/AGENT INFORMATION:
     NAME: Clough, David W.
     REGISTRATION NUMBER: 36,107
     REFERENCE/DOCKET NUMBER: 01017/34451
   TELECOMMUNICATION INFORMATION:
     TELEPHONE: (312) 474-6300
     TELEFAX: (312) 474-0448
  INFORMATION FOR SEQ ID NO: 3:
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;
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; Patent No. 6512164
; GENERAL INFORMATION:
 APPLICANT: Famodu, Omolayo O.
 APPLICANT: Rafalski, J. Antoni
 TITLE OF INVENTION: Thiamine Biosynthetic Enzymes
; FILE REFERENCE: BB1372 US NA
; CURRENT APPLICATION NUMBER: US/09/594,506
 CURRENT FILING DATE: 2000-06-15
; PRIOR APPLICATION NUMBER: 60/139,556
 PRIOR FILING DATE: 1999-06-16
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  ORGANISM: Triticum aestivum
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; Patent No. 6436657
; GENERAL INFORMATION:
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; APPLICANT: Morakinyo, Layo O.
 APPLICANT: Orozco Jr, Emil M.
  TITLE OF INVENTION: TETRAHYDROFOLATE METABOLIC ENZYMES
; FILE REFERENCE: BB1322 US NA
; CURRENT APPLICATION NUMBER: US/09/465,558
; CURRENT FILING DATE: 1999-12-17
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; EARLIER FILING DATE: 1998-12-18
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  TYPE: DNA
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; Patent No. 5789222
; GENERAL INFORMATION:
   APPLICANT: KELLY, ROSEMARIE
   APPLICANT: REGISTER, ELIZABETH A APPLICANT: MASUREKAR, PRAKASH S
   TITLE OF INVENTION: P5C REDUCTASE GENE FROM ZALERION
   TITLE OF INVENTION: ARBORICOLA
   NUMBER OF SEQUENCES: 2
   CORRESPONDENCE ADDRESS:
     ADDRESSEE: MERCK & CO., INC.
   STREET: 126 E. LINCOLN AVENUE
     CITY: RAHWAY
     STATE: NEW JERSEY
     COUNTRY: US
     ZIP: 07065-0900
   COMPUTER READABLE FORM:
     MEDIUM TYPE: Floppy disk
     COMPUTER: IBM PC compatible
     OPERATING SYSTEM: PC-DOS/MS-DOS
     SOFTWARE: PatentIn Release #1.0, Version #1.30
  CURRENT APPLICATION DATA:
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     FILING DATE: 23-JUN-1995
     CLASSIFICATION:
   ATTORNEY/AGENT INFORMATION:
     NAME: KORSEN, ELLIOTT
     REGISTRATION NUMBER: 32,705
     REFERENCE/DOCKET NUMBER: 19453PV
   TELECOMMUNICATION INFORMATION:
     TELEPHONE: 908-594-5493
     TELEFAX: 908-594-4720
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     TOPOLOGY: linear
    MOLECULE TYPE: cDNA
    HYPOTHETICAL: NO
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    FEATURE:
     NAME/KEY: CDS
     LOCATION: 47..960
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; Sequence 22, Application US/09182816
; Patent No. 6143542
; GENERAL INFORMATION:
; APPLICANT: Wisnewski, Nancy
; APPLICANT: Silver, Garv M.
 APPLICANT: Lo, Katherine C.
  APPLICANT: Brandt, Kevin S.
  TITLE OF INVENTION: NOVEL FLEA EPOXIDE HYDROLASE NUCLEIC ACID MOLECULES,
  TITLE OF INVENTION: PROTEINS AND USES THEREOF
  FILE REFERENCE: FC-3-C1
 CURRENT APPLICATION NUMBER: US/09/182,816
  CURRENT FILING DATE: 1998-10-29
  EARLIER APPLICATION NUMBER: 08/989,510
 EARLIER FILING DATE: 1997-12-12
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; SEQ ID NO 22
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; Patent No. 6143542
; GENERAL INFORMATION:
; APPLICANT: Wisnewski, Nancy
; APPLICANT: Silver, Gary M.
 APPLICANT: Lo, Katherine C.
  APPLICANT: Brandt, Kevin S.
  TITLE OF INVENTION: NOVEL FLEA EPOXIDE HYDROLASE NUCLEIC ACID MOLECULES,
  TITLE OF INVENTION: PROTEINS AND USES THEREOF
; FILE REFERENCE: FC-3-C1
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  CURRENT FILING DATE: 1998-10-29
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  EARLIER FILING DATE: 1997-12-12
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; GENERAL INFORMATION:
 APPLICANT: Wisnewski, Nancy
 APPLICANT: Silver, Gary M.
 APPLICANT: Lo, Katherine C.
 APPLICANT: Brandt, Kevin S.
 TITLE OF INVENTION: FLEA EPOXIDE HYDROLASE PROTEINS AND USES THEREOF
; FILE REFERENCE: FC-3-C1-1
; CURRENT APPLICATION NUMBER: US/09/471,528
 CURRENT FILING DATE: 1999-12-27
; EARLIER APPLICATION NUMBER: 09/182,816
; EARLIER FILING DATE: 1998-10-29
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 EARLIER FILING DATE: 1997-12-12
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; Patent No. 6153397

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; GENERAL INFORMATION:
 APPLICANT: Wisnewski, Nancy
 APPLICANT: Silver, Gary M.
 APPLICANT: Lo, Katherine C.
; APPLICANT: Brandt, Kevin S.
 TITLE OF INVENTION: FLEA EPOXIDE HYDROLASE PROTEINS AND USES THEREOF
; FILE REFERENCE: FC-3-C1-1
  CURRENT APPLICATION NUMBER: US/09/471,528
  CURRENT FILING DATE: 1999-12-27
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  EARLIER FILING DATE: 1997-12-12
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; GENERAL INFORMATION:
; APPLICANT: Wisnewski, Nancy
 APPLICANT: Silver, Gary M.
 APPLICANT: Lo, Katherine C.
 APPLICANT: Brandt, Kevin S.
  TITLE OF INVENTION: FLEA EPOXIDE HYDROLASE PROTEINS AND USES THEREOF
  FILE REFERENCE: FC-3-C1-1
  CURRENT APPLICATION NUMBER: US/09/634,530
  CURRENT FILING DATE: 2000-08-08
  PRIOR APPLICATION NUMBER: 09/471,528
  PRIOR FILING DATE: 1999-12-27
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  PRIOR FILING DATE: 1998-10-29
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GenCore version 5.1.6 Copyright (c) 1993 - 2004 Compugen Ltd.

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Title: US-10-056-884A-1

Perfect score: 3468

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SUMMARIES

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	15	79	2.3	1492	9	US-09-925-299-112	Sequence 112, App
	16	79	2.3	1492	11	US-09-925-299-112	Sequence 112, App
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ALIGNMENTS

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US-10-056-884-1
; Sequence 1, Application US/10056884
; Publication No. US20030032786A1
; GENERAL INFORMATION:
  APPLICANT: Bristol-Myers Squibb Company
  TITLE OF INVENTION: POLYNUCLEOTIDE ENCODING A NOVEL HUMAN POTASSIUM CHANNEL
BETA-SUBUNIT,
  TITLE OF INVENTION: K+betaM2
  FILE REFERENCE: D0076 NP
  CURRENT APPLICATION NUMBER: US/10/056,884
  CURRENT FILING DATE: 2002-01-24
   PRIOR APPLICATION NUMBER: US 60/263,872
  PRIOR FILING DATE: 2001-01-24
   PRIOR APPLICATION NUMBER: US 60/269,794
   PRIOR FILING DATE: 2001-02-14
  NUMBER OF SEQ ID NOS: 73
   SOFTWARE: PatentIn version 3.0
; SEQ ID NO 1
   LENGTH: 3468
    TYPE: DNA
;
    ORGANISM: Homo sapiens
    FEATURE:
    NAME/KEY: CDS
    LOCATION: (515)..(1798)
US-10-056-884-1
  Query Match
                          100.0%; Score 3468; DB 15; Length 3468;
  Best Local Similarity
                          100.0%; Pred. No. 0;
 Matches 3468; Conservative
                                 0; Mismatches
                                                   0; Indels
                                                                     Gaps
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; GENERAL INFORMATION:
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BETA-SUBUNIT,
  TITLE OF INVENTION: K+betaM2
  FILE REFERENCE: D0076 NP
  CURRENT APPLICATION NUMBER: US/10/056,884
  CURRENT FILING DATE: 2002-01-24
  PRIOR APPLICATION NUMBER: US 60/263,872
  PRIOR FILING DATE: 2001-01-24
  PRIOR APPLICATION NUMBER: US 60/269,794
  PRIOR FILING DATE: 2001-02-14
  NUMBER OF SEQ ID NOS: 73
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US-10-056-884-3
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US-10-029-386-10927/c
; Sequence 10927, Application US/10029386
; Publication No. US20030194704A1
; GENERAL INFORMATION:
  APPLICANT: Penn, Sharron G.
; APPLICANT: Rank, David R.
; APPLICANT: Hanzel, David K.
  TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES
USEFUL FOR GENE
  TITLE OF INVENTION: EXPRESSION ANALYSIS TWO
  FILE REFERENCE: AEOMICA-X-2
  CURRENT APPLICATION NUMBER: US/10/029,386
  CURRENT FILING DATE: 2001-12-20
  NUMBER OF SEQ ID NOS: 34288
  SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
; SEQ ID NO 10927
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   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
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   OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 0.44
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US-10-029-386-10927
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US-10-029-386-24630/c
; Sequence 24630, Application US/10029386
; Publication No. US20030194704A1
; GENERAL INFORMATION:
 APPLICANT: Penn, Sharron G.
 APPLICANT: Rank, David R.
 APPLICANT: Hanzel, David K.
  TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES
USEFUL FOR GENE
  TITLE OF INVENTION: EXPRESSION ANALYSIS TWO
  FILE REFERENCE: AEOMICA-X-2
  CURRENT APPLICATION NUMBER: US/10/029,386
  CURRENT FILING DATE: 2001-12-20
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   ORGANISM: Homo sapiens
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   OTHER INFORMATION: MAP TO AC008716.6
   OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 0.44
   OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 0.55
   OTHER INFORMATION: SWISSPROT HIT: P19836, EVALUE 2.30e+00
   OTHER INFORMATION: NT HIT: AB037738.1, EVALUE 0.00e+00
US-10-029-386-24630
                       6.9%; Score 240; DB 13; Length 279;
 Query Match
 Best Local Similarity
                      98.0%; Pred. No. 1.4e-46;
 Matches 243; Conservative
                          0; Mismatches
                                            5; Indels
       1336 TGTCTTCTACCGTGAGCCTTCCAGATGGTCACCCTCACACTGCGATTGCTGCTGCAAGAA 1395
Qу
```

```
248 TTTCTTTTCAGGTGAGCCTTCCAGATGGTCACCCTCACACTGCGATTGCTGCTGCAAGAA 189
Db
       1396 TGGCAAAGGTGACAAAGAAGGGGAGAGCGGCACGTCTTGCAATGACCTCTCCACATCTAG 1455
Qу
           188 TGGCAAAGGTGACAAAGAAGGGGAGAGCGGCACGTCTTGCAATGACCTCTCCACATCTAG 129
Db
       1456 CTGCGACAGCCAGTCTGAGGCCAGCTCTCCCCAGGAGACGGTCATCTGTGGTCCCGTGAC 1515
Qу
            Db
        128 CTGCGACAGCCAGTCTGAGGCCAGCTCTCCCCAGGAGACGGTCATCTGTGGTCCCGTGAC 69
       1516 ACGCCAGACCAACATCCAGACTCTGGACCGTCCCATCAAGAAGGGCCCTGTCCAGCTGAT 1575
Qу
           68 ACGCCAGACCAACATCCAGACTCTGGACCGTCCCATCAAGAAGGGCCCTGTCCAGCTGAT 9
Db
       1576 CCAACAGT 1583
Qу
           11111111
          8 CCAACAGT 1
Db
RESULT 5
US-10-060-036-4467
; Sequence 4467, Application US/10060036
; Publication No. US20030073144A1
; GENERAL INFORMATION:
  APPLICANT: Benson, Darin R.
 APPLICANT: Kalos, Michael D.
  APPLICANT: Lodes, Michael J.
  APPLICANT: Persing, David H.
 APPLICANT: Hepler, William T.
  APPLICANT: Jiang, Yuqiu
  TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY
  TITLE OF INVENTION: AND DIAGNOSIS OF PANCREATIC CANCER
 FILE REFERENCE: 210121.566
; CURRENT APPLICATION NUMBER: US/10/060,036
  CURRENT FILING DATE: 2002-01-30
  NUMBER OF SEQ ID NOS: 4560
  SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 4467
   LENGTH: 632
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: misc feature
   LOCATION: 552, 569
   OTHER INFORMATION: n = A, T, C or G
US-10-060-036-4467
                      5.9%; Score 205; DB 15; Length 632;
 Query Match
 Best Local Similarity 93.4%; Pred. No. 4.4e-38;
                           0; Mismatches 15; Indels
 Matches 214; Conservative
                                                       0; Gaps
Qу
       3206 AGGTACCAATAGCTCTTTCATAGACTTGTGCTACAAGAAGGTTAAAAGACCAGTTTTATT 3265
            Db
         15 AGGTACCAATAGCTCTTTCATAGACTTGTGCTACAAGAAGGTTAAAAGACCAGTTTTATT 74
       3266 TTCAGCATTCCTCATGCATTTCAGTGGTAACCAAAAAATAATTTGTCAATTAATAGTTGT 3325
Qу
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75 TTCAGCATTCCTCATGCATTTCAGTGGTAACCAAAAAATAATTTGTCAATTAATAGTTGT 134
Db
     Qy
         Db
     Qу
                           Db
      195 GGTAATAAAGGCAATTGGATGATATCTGTAGGAGGAAAACAATGACTAA 243
RESULT 6
US-10-060-036-564
; Sequence 564, Application US/10060036
; Publication No. US20030073144A1
; GENERAL INFORMATION:
 APPLICANT: Benson, Darin R.
 APPLICANT: Kalos, Michael D.
 APPLICANT: Lodes, Michael J.
 APPLICANT: Persing, David H.
 APPLICANT: Hepler, William T.
 APPLICANT: Jiang, Yuqiu
 TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY
  TITLE OF INVENTION: AND DIAGNOSIS OF PANCREATIC CANCER
  FILE REFERENCE: 210121.566
 CURRENT APPLICATION NUMBER: US/10/060,036
 CURRENT FILING DATE: 2002-01-30
 NUMBER OF SEQ ID NOS: 4560
 SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 564
  LENGTH: 614
  TYPE: DNA
  ORGANISM: Homo sapiens
  FEATURE:
  NAME/KEY: misc feature
  LOCATION: 534, 551, 575, 576
  OTHER INFORMATION: n = A, T, C or G
US-10-060-036-564
 Query Match
                 5.8%; Score 201; DB 15; Length 614;
 Best Local Similarity 93.3%; Pred. No. 3.9e-37;
 Matches 210; Conservative
                     0; Mismatches 15; Indels
                                          0; Gaps
                                                  0;
     3210 ACCAATAGCTCTTTCATAGACTTGTGCTACAAGAAGGTTAAAAGACCAGTTTTATTTTCA 3269
Qy
         Db
       1 ACCAATAGCTCTTTCATAGACTTGTGCTACAAGAAGGTTAAAAGACCAGTTTTATTTTCA 60
Qу
     3270 GCATTCCTCATGCATTTCAGTGGTAACCAAAAATAATTTGTCAATTAATAGTTGTGTGC 3329
         Db
       61 GCATTCCTCATGCATTTCAGTGGTAACCAAAAAATAATTTGTCAATTAATAGTTGTGTGC 120
Qу
     Db
      Qу
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RESULT 7
US-10-080-980-1
; Sequence 1, Application US/10080980
; Publication No. US20030036115A1
; GENERAL INFORMATION:
  APPLICANT: Bristol-Myers Squibb Company
  TITLE OF INVENTION: POLYNUCLEOTIDE ENCODING A NOVEL HUMAN POTASSIUM CHANNEL
BETA-SUBUNIT,
  TITLE OF INVENTION: K+betaM6, EXPRESSED HIGHLY IN THE SMALL INTESTINE
  FILE REFERENCE: D0121 NP
  CURRENT APPLICATION NUMBER: US/10/080,980
  CURRENT FILING DATE: 2002-02-21
  PRIOR APPLICATION NUMBER: US 60/270,132
  PRIOR FILING DATE: 2001-02-21
  PRIOR APPLICATION NUMBER: US 60/278,953
  PRIOR FILING DATE: 2001-03-27
  NUMBER OF SEQ ID NOS: 74
  SOFTWARE: PatentIn version 3.0
; SEQ ID NO 1
   LENGTH: 2052
   TYPE: DNA
   ORGANISM: homo sapiens
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (121)..(1095)
US-10-080-980-1
 Query Match
                       4.8%; Score 167; DB 15; Length 2052;
 Best Local Similarity
                      64.6%; Pred. No. 9.4e-29;
 Matches 267: Conservative
                            0; Mismatches 140; Indels
                                                       6; Gaps
                                                                  1;
        967 CCCCCCTTCCTCCCTGCCGCCGCCGCCGCGGGTTTCATTACTGTGGGTTACAG 1026
Qy
                        111
                              1 1 111
            705 CACGCCGTCCCAGTCGCTGGACGGCAGCCGGCGCTCGGGCTACATCACCATCGGCTACCG 764
Db
       1027 AGGATCCTGCACCTTGGGCAGAGAGGGGACAGGCAGATGCCAAGTTTCGGAGAGTTCCCCG 1086
Qу
            765 CGGCTCCTACACCATCGGCGGGGCGGACGCGAGGCGACGCCAAGTTCCGGCGAGTGGCGCG 824
Db
       1087 GATTTTGGTTTGTGGAAGGATTTCCTTGGCAAAAGAAGTCTTTGGAGAAACTTTGAATGA 1146
QУ
                  825 CATCACCGTTTGCGGAAAGACGTCGCTGGCCAAGGAGGTGTTTGGGGACACCCTGAACGA 884
Db
Qу
       1147 AAGCAGAGACCCTGATCGAGCCCCAGAAAGATACACCTCCAGATTTTATCTCAAATTCAA 1206
            885 AAGCCGGGACCCCGACCGTCCCCGGAGCGCTACACCTCGCGCTATTACCTCAAGTTCAA 944
Db
Qy
       1207 GCACCTGGAAAGGGCTTTTGATATGTTGTCAGAGTGTGGATTCCACATGGTGGCCTGTAA 1266
                      Db
        945 CTTCCTGGAGCAGGCCTTCGACAAGCTGTCCGAGTCGGGCTTCCACATGGTGGCGTGCAG 1004
       1267 CTCATCGGTGACAGCATCTTT-----CATCAACCAATATACAGATGACAAGATCTGGTC 1320
Qу
            111 111 11 111
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1005 CTCCACGGGCACCTGCGCCTTTGCCAGCAGCACCAGAGCGAGGAGAAGATCTGGAC 1064
Db
        1321 AAGCTACACTGAATATGTCTTCTACCGTGAGCCTTCCAGATGGTCACCCTCAC 1373
Οv
              1065 CAGCTACACCGAGTACGTCTTCTGCAGGGAGTGAGCTCCCCAGACCCCCTCGC 1117
Db
RESULT 8
US-10-029-386-20178/c
; Sequence 20178, Application US/10029386
; Publication No. US20030194704A1
; GENERAL INFORMATION:
  APPLICANT: Penn, Sharron G.
  APPLICANT: Rank, David R.
  APPLICANT: Hanzel, David K.
  TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES
USEFUL FOR GENE
  TITLE OF INVENTION: EXPRESSION ANALYSIS TWO
  FILE REFERENCE: AEOMICA-X-2
  CURRENT APPLICATION NUMBER: US/10/029,386
  CURRENT FILING DATE: 2001-12-20
  NUMBER OF SEQ ID NOS: 34288
  SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
; SEO ID NO 20178
   LENGTH: 978
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   OTHER INFORMATION: MAP TO AC000403.1
   OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 12
   OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 12
   OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 12
   OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 15
   OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 13
   OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 7.7
   OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 8.1
   OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 6.3
   OTHER INFORMATION: SWISSPROT HIT: Q14681, EVALUE 2.00e-04
   OTHER INFORMATION: EST HUMAN HIT: BG387727.1, EVALUE 8.00e-64
   OTHER INFORMATION: NT HIT: gi16163086, EVALUE 0.00e+00
US-10-029-386-20178
 Query Match
                         4.5%; Score 156.2; DB 13; Length 978;
 Best Local Similarity 65.6%; Pred. No. 2.2e-26;
 Matches 246; Conservative
                             0; Mismatches 123; Indels
                                                            6; Gaps
                                                                       1:
         967 CCCCCCTTCCTCCCTGCCGCCGCCGCACGCGCAGTGGGGTTTCATTACTGTGGGTTACAG 1026
Qу
             1 1 11 111 111
                                Db
         376 CACGCCGTCCCAGTCGCTGGACGGCAGCCGGCGCTCGGGCTACATCACCATCGGCTACCG 317
        1027 AGGATCCTGCACCTTGGGCAGAGAGGGGACAGGCAGATGCCAAGTTTCGGAGAGTTCCCCG 1086
Qу
              Db
         316 CGGCTCCTACACCATCGGGCGGGACGCGCAGGCGGACGCCAAGTTCCGGCGAGTGGCGCG 257
        1087 GATTTTGGTTTGTGGAAGGATTTCCTTGGCAAAAGAAGTCTTTGGAGAAACTTTGAATGA 1146
Qy
                   11111 1111 11 11 11 1111 11 11 11 1111 11 11 11
              11
Dh
         256 CATCACCGTTTGCGGAAAGACGTCGCTGGCCAAGGAGGTGTTTGGGGACACCCTGAACGA 197
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1147 AAGCAGAGCCCTGATCGAGCCCCAGAAAGATACACCTCCAGATTTTATCTCAAATTCAA 1206
Qy
            196 AAGCCGGGACCCCGACCGTCCCCCGGAGCGCTACACCTCGCGCTATTACCTCAAGTTCAA 137
Db
Qv
        1207 GCACCTGGAAAGGGCTTTTGATATGTTGTCAGAGTGTGGATTCCACATGGTGGCCTGTAA 1266
                      136 CTTCCTGGAGCAGGCCTTCGACAAGCTGTCCGAGTCGGGCTTCCACATGGTGGCGTGCAG 77
Dh
        1267 CTCATCGGTGACAGCATCTTT-----CATCAACCAATATACAGATGACAAGATCTGGTC 1320
Qу
                          1 11
                                    Db
         76 CTCCACGGGCACCTGCGCCTTTGCCAGCAGCACCAGAGCGAGGAGGACAAGATCTGGAC 17
Qу
       1321 AAGCTACACTGAATA 1335
             11111111111
Db
         16 CAGCTACACCGAGTA 2
RESULT 9
US-10-080-980-8
; Sequence 8, Application US/10080980
; Publication No. US20030036115A1
; GENERAL INFORMATION:
  APPLICANT: Bristol-Myers Squibb Company
  TITLE OF INVENTION: POLYNUCLEOTIDE ENCODING A NOVEL HUMAN POTASSIUM CHANNEL
BETA-SUBUNIT,
  TITLE OF INVENTION: K+betaM6, EXPRESSED HIGHLY IN THE SMALL INTESTINE
  FILE REFERENCE: D0121 NP
  CURRENT APPLICATION NUMBER: US/10/080,980
  CURRENT FILING DATE: 2002-02-21
  PRIOR APPLICATION NUMBER: US 60/270,132
  PRIOR FILING DATE: 2001-02-21
  PRIOR APPLICATION NUMBER: US 60/278,953
  PRIOR FILING DATE: 2001-03-27
  NUMBER OF SEQ ID NOS: 74
  SOFTWARE: PatentIn version 3.0
; SEQ ID NO 8
   LENGTH: 688
   TYPE: DNA
   ORGANISM: homo sapiens
   FEATURE:
   NAME/KEY: misc feature
   OTHER INFORMATION: wherein "N" is equal to "A", "C", "G" or "T".
US-10-080-980-8
                       3.0%; Score 104.6; DB 15; Length 688;
 Query Match
                       51.1%; Pred. No. 3e-14;
 Best Local Similarity
 Matches 192; Conservative
                            0; Mismatches 178; Indels
                                                         6; Gaps
                                                                    1;
QУ
        563 TCCGCAGTTCCCAACTCCTTCCCTGAGGTGGTAGAGCTGAATGTCGGGGGGTCAAGTTTAT 622
            186 TCCGCGGAGCCACCGCTCTTCCCCGACATCGTGGAGCTGAACGTGGGGGGCCCAGGTGTAC 245
Qу
        623 TTTACTCGCCATTCCACATTGATAAGCATCCCTCATTCCCTCTGTGGAAAATGTTTTCC 682
             1 11 11 1 1 1 1 1 1 1
                                    11111 1
Db
        246 GTGACCCGGCGCTGCACGGTGGTGTCGGTGCCCGACTCGCTGCTCTGGCGCATGTTCACG 305
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683 CCAAAGAGAGACACGGCTAATGATCTAGCCAAGGACTCCAAGGGAAGGTTTTTCATTGAC 742
Qу
          Db
       Qу
           360 CGGGACGGCTTCCTCTTCCGCTACATCCTGGATTACCTGCGGGACTTGCAGCTCGTGCTG 419
Db
       803 CCTGATCACTTTCCAGAAAAAGGAAGACTGAAAAGGGAAGCTGAATACTTCCAGCTCCCA 862
Qу
           Db
       420 CCCGACTACTTCCCCGAGCGCAGCCGGCTGCAGCGCGAGGCCGAGTACTTCGAGCTGCCA 479
       863 GACTTGGTCAAACTCCTGACCCCGATGAAATCAAGCAAAGCCCAGATGAATTCTGCCAC 922
Qу
Db
       923 AGTGACTTTGAAGATG 938
Qу
                 1 1 1
Db
       540 NNNNNNTGCACAAGG 555
RESULT 10
US-09-918-995-2311
; Sequence 2311, Application US/09918995
; Publication No. US20030073623A1
; GENERAL INFORMATION:
  APPLICANT: Hyseq, Inc.
  TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED
  TITLE OF INVENTION: FROM VARIOUS cDNA LIBRARIES
  FILE REFERENCE: 20411-756
  CURRENT APPLICATION NUMBER: US/09/918,995
  CURRENT FILING DATE: 2001-07-30
  PRIOR APPLICATION NUMBER: US/09/235,076
  PRIOR FILING DATE: 1999-01-20
  NUMBER OF SEQ ID NOS: 38054
  SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 2311
   LENGTH: 249
   TYPE: DNA
   ORGANISM: Homo sapiens
US-09-918-995-2311
 Query Match
                    2.5%; Score 87; DB 11; Length 249;
 Best Local Similarity 65.2%; Pred. No. 2.4e-10;
 Matches 161; Conservative
                         0; Mismatches 80; Indels
                                                  6; Gaps
                                                           2;
Qу
       1236 CAGAGTGTGGATTCCACATGGTGGCCTGTAACTCATCGGTGACAGCATCTTTCATCAACC 1295
                 Db
         1 CCGAGGCCGCTTCCACATGGTGGCGTGTAACTCCTCGGGCACCGCCGCCTTCGTCAACC 60
Qу
       1296 AATATACAGATGACAAGATCTGGTCAAGCTACACTGAATATGTCTTCTACCGTGAGCCT- 1354
                 11 111111111
                              61 AGTACCGCGACGACAAGATCTGGAGCAGCTACACCGAGTACATTTTCTTCCGACCACCTC 120
Db
Qу
       1355 --TCCAGATGGTCACCCTCACACTGCGATTGCTGCTGCAAGAATGGCAAAG---GTGACA 1409
              1 1
                   ++++
                         111
                               1 1
                                          Db
       121 AGAAAATAGTATCACCTAAACAAGAACATGAAGATAGGATACATGACCAAGTCACTGATA 180
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1410 AAGAAGGGGAGAGCGCACGTCTTGCAATGACCTCTCCACATCTAGCTGCGACAGCCAGT 1469
Qу
              Db
         181 AAGGAAGTGAAAGTGGGACTTCCTGGAATGAGCTCTTCACTTCCAGTTGGGACAGCCATT 240
Qу
        1470 CTGAGGC 1476
             1 11111
Db
         241 CAGAGGC 247
RESULT 11
US-09-814-353-4862/c
; Sequence 4862, Application US/09814353
; Publication No. US20030165831A1
; GENERAL INFORMATION:
  APPLICANT: Lee, John
  APPLICANT: Thompson, Pamela
  APPLICANT: Lillie, James
  TITLE OF INVENTION: NOVEL GENES, COMPOSITIONS, KITS, AND METHODS FOR
  TITLE OF INVENTION: IDENTIFICATION, ASSESSMENT, PREVENTION, AND
  TITLE OF INVENTION: THERAPY OF OVARIAN CANCER
  FILE REFERENCE: MRI-006B
  CURRENT APPLICATION NUMBER: US/09/814,353
  CURRENT FILING DATE: 2001-03-21
  PRIOR APPLICATION NUMBER: US 60/191,031
  PRIOR FILING DATE: 2000-03-21
  PRIOR APPLICATION NUMBER: US 60/207,124
  PRIOR FILING DATE: 2000-05-25
  PRIOR APPLICATION NUMBER: US 60/211,940
  PRIOR FILING DATE: 2000-06-15
  PRIOR APPLICATION NUMBER: US 60/216,820
  PRIOR FILING DATE: 2000-07-07
  PRIOR APPLICATION NUMBER: US 60/220,661
  PRIOR FILING DATE: 2000-07-25
  PRIOR APPLICATION NUMBER: US 60/257,672
  PRIOR FILING DATE: 2000-12-21
  NUMBER OF SEQ ID NOS: 22037
  SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 4862
   LENGTH: 496
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: misc feature
   LOCATION: 156, 157, 160, 161, 162, 163, 164, 165, 167, 168, 169, 170,
   LOCATION: 171, 172, 173, 174, 175, 196, 197, 198, 200, 203, 205, 206,
   LOCATION: 219, 220, 228, 232, 240, 241, 244, 245, 246, 247, 249, 250,
   LOCATION: 252, 253, 256, 258, 260, 263, 264, 265, 267, 268, 269
   OTHER INFORMATION: n = A, T, C or G
;
   FEATURE:
   NAME/KEY: misc feature
    LOCATION: 270, 271, 274, 275, 280, 287, 288, 289, 290, 303, 306, 317,
    LOCATION: 318, 322, 331, 347, 348, 355, 361, 362, 364, 367, 368, 369,
    LOCATION: 381, 383, 388, 393, 398, 404, 408, 409, 410, 411, 412, 413,
    LOCATION: 414, 415, 416, 417, 418, 419, 420, 421, 423, 424, 435
   OTHER INFORMATION: n = A, T, C or G
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FEATURE:

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NAME/KEY: misc feature
  LOCATION: 436, 438, 441, 450, 451, 452, 453, 454, 455, 456, 457, 458,
  LOCATION: 470, 471, 472, 475, 477, 478, 481, 482, 484
  OTHER INFORMATION: n = A, T, C or G
US-09-814-353-4862
                    2.3%; Score 80.6; DB 13; Length 496;
 Ouerv Match
 Best Local Similarity 47.3%; Pred. No. 1.2e-08;
 Matches 131; Conservative 0; Mismatches 146; Indels
                                                0; Gaps
                                                           0;
      3192 TCTGTATTTTACTAAGGTACCAATAGCTCTTTCATAGACTTGTGCTACAAGAAGGTTAAA 3251
Qу
                      1 | 1 | 1 | 1
Db
       343 TTTTTTTTTCCNCCCCTTTCNTTTNNAATTAAAAAANATNTTTTTTCCCCAANNNNAAA 284
      3252 AGACCAGTTTTATTTTCAGCATTCCTCATGCATTTCAGTGGTAACCAAAAAATAATTTGT 3311
Qу
          1 | | |
                         11
                              Db
       283 AAANAAAANNAANNNNTTNNTTNANTNTTNNTNNCNNNNGGNNAAAAAAANTTTNTTTT 224
      Qy
              - 1 1
Db
       Qу
             Db
       Qу
      3432 AAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA 3468
          RESULT 12
US-09-814-353-11159/c
; Sequence 11159, Application US/09814353
; Publication No. US20030165831A1
; GENERAL INFORMATION:
; APPLICANT: Lee, John
; APPLICANT: Thompson, Pamela
  APPLICANT: Lillie, James
  TITLE OF INVENTION: NOVEL GENES, COMPOSITIONS, KITS, AND METHODS FOR
  TITLE OF INVENTION: IDENTIFICATION, ASSESSMENT, PREVENTION, AND TITLE OF INVENTION: THERAPY OF OVARIAN CANCER
  FILE REFERENCE: MRI-006B
  CURRENT APPLICATION NUMBER: US/09/814,353
  CURRENT FILING DATE: 2001-03-21
  PRIOR APPLICATION NUMBER: US 60/191,031
  PRIOR FILING DATE: 2000-03-21
  PRIOR APPLICATION NUMBER: US 60/207,124
  PRIOR FILING DATE: 2000-05-25
  PRIOR APPLICATION NUMBER: US 60/211,940
  PRIOR FILING DATE: 2000-06-15
  PRIOR APPLICATION NUMBER: US 60/216,820
  PRIOR FILING DATE: 2000-07-07
  PRIOR APPLICATION NUMBER: US 60/220,661
  PRIOR FILING DATE: 2000-07-25
; PRIOR APPLICATION NUMBER: US 60/257,672
; PRIOR FILING DATE: 2000-12-21
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NUMBER OF SEQ ID NOS: 22037
  SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 11159
   LENGTH: 496
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: misc feature
   LOCATION: 156, 157, 160, 161, 162, 163, 164, 165, 167, 168, 169, 170,
   LOCATION: 171, 172, 173, 174, 175, 196, 197, 198, 200, 203, 205, 206,
   LOCATION: 219, 220, 228, 232, 240, 241, 244, 245, 246, 247, 249, 250,
   LOCATION: 252, 253, 256, 258, 260, 263, 264, 265, 267, 268, 269
   OTHER INFORMATION: n = A, T, C or G
   FEATURE:
   NAME/KEY: misc feature
   LOCATION: 270, 271, 274, 275, 280, 287, 288, 289, 290, 303, 306, 317,
  LOCATION: 318, 322, 331, 347, 348, 355, 361, 362, 364, 367, 368, 369,
   LOCATION: 381, 383, 388, 393, 398, 404, 408, 409, 410, 411, 412, 413,
   LOCATION: 414, 415, 416, 417, 418, 419, 420, 421, 423, 424, 435
   OTHER INFORMATION: n = A, T, C or G
   FEATURE:
   NAME/KEY: misc feature
   LOCATION: 436, 438, 441, 450, 451, 452, 453, 454, 455, 456, 457, 458,
   LOCATION: 470, 471, 472, 475, 477, 478, 481, 482, 484
   OTHER INFORMATION: n = A, T, C or G
US-09-814-353-11159
 Query Match
                    2.3%; Score 80.6; DB 13; Length 496;
 Best Local Similarity
                    47.3%; Pred. No. 1.2e-08;
 Matches 131; Conservative
                         0; Mismatches 146;
                                          Indels
                                                  0; Gaps
                                                            0:
       3192 TCTGTATTTTACTAAGGTACCAATAGCTCTTTCATAGACTTGTGCTACAAGAAGGTTAAA 3251
Qу
                                11 1 1
           11
Db
       343 TTTTTTTTTCCNCCCCTTTCNTTTNNAATTAAAAAANATNTTTTTTCCCCAANNNNAAA 284
       3252 AGACCAGTTTTATTTTCAGCATTCCTCATGCATTTCAGTGGTAACCAAAAAATAATTTGT 3311
Qу
          - 1
                                              Db
       283 AAANAAAANNAANNNNTTNANTTNANTNTTNNTNNCNNNNGGNNAAAAAAANTTTNTTTT 224
Qу
       1 1
Db
       Qy
       1
                   Db
       Qy
       3432 AAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA3468
           Db
       103 АААААААААААААААААААААААААААААААААА
RESULT 13
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US-10-056-884-8/c

[;] Sequence 8, Application US/10056884

[;] Publication No. US20030032786A1

[;] GENERAL INFORMATION:

```
; APPLICANT: Bristol-Myers Squibb Company
  TITLE OF INVENTION: POLYNUCLEOTIDE ENCODING A NOVEL HUMAN POTASSIUM CHANNEL
BETA-SUBUNIT,
; TITLE OF INVENTION: K+betaM2
; FILE REFERENCE: D0076 NP
; CURRENT APPLICATION NUMBER: US/10/056,884
; CURRENT FILING DATE: 2002-01-24
; PRIOR APPLICATION NUMBER: US 60/263,872
 PRIOR FILING DATE: 2001-01-24
  PRIOR APPLICATION NUMBER: US 60/269,794
 PRIOR FILING DATE: 2001-02-14
; NUMBER OF SEQ ID NOS: 73
  SOFTWARE: PatentIn version 3.0
; SEO ID NO 8
   LENGTH: 80
   TYPE: DNA
   ORGANISM: Artificial Sequence
   FEATURE:
   OTHER INFORMATION: Synthetic Oligonucleotide Modified To Contain Biotin at
the 5 Pr
   OTHER INFORMATION: ime En
US-10-056-884-8
 Query Match 2.3%; Score 80; DB 15; Length 80; Best Local Similarity 100.0%; Pred. No. 5.4e-09;
 Matches 80; Conservative 0; Mismatches 0; Indels
                                                               0; Gaps
                                                                           0:
Qy
         783 GGGACAGGCAGGTGGTCCTGCCTGATCACTTTCCAGAAAAGGAAGACTGAAAAGGGAAG 842
             Db
          80 GGGACAGGCAGGTGGTCCTGATCACTTTCCAGAAAAAGGAAGACTGAAAAGGGAAG 21
         843 CTGAATACTTCCAGCTCCCA 862
Qу
             111111111111111111111
Db
          20 CTGAATACTTCCAGCTCCCA 1
RESULT 14
US-09-834-975-451/c
; Sequence 451, Application US/09834975
; Patent No. US20020110815A1
; GENERAL INFORMATION:
; APPLICANT: Lillie, James
; APPLICANT: Brown, Jeffrey
; APPLICANT: Bolt, Andrew
; APPLICANT: Van Huffel, Christophe
  TITLE OF INVENTION: NOVEL GENES, COMPOSITIONS AND METHODS
  TITLE OF INVENTION: FOR THE IDENTIFICATION, ASSESSMENT, PREVENTION, AND
THERAPY
; TITLE OF INVENTION: OF HUMAN CANCERS
; FILE REFERENCE: MRI-016B
; CURRENT APPLICATION NUMBER: US/09/834,975
; CURRENT FILING DATE: 2001-04-13
; PRIOR APPLICATION NUMBER: 60/197,538
; PRIOR FILING DATE: 2000-04-14
; NUMBER OF SEQ ID NOS: 1046
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 451
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  OTHER INFORMATION: n = A, T, C or G
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         Db
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Qу
         Db
      Qу
     3287 CAGTGGTAACCAAAAAATAATTTGTCAATTAATAGTTGTGCCAAGCACTCCTAATTTG 3346
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     Qу
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     Qу
        Db
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Qу
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      21 AA 20
Db
RESULT 15
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; Sequence 112, Application US/09925299
; Patent No. US20020055627A1
; GENERAL INFORMATION:
 APPLICANT: Rosen et al.
 TITLE OF INVENTION: Nucleic Acids, Proteins and Antibodies
 FILE REFERENCE: PA102
 CURRENT APPLICATION NUMBER: US/09/925,299
 CURRENT FILING DATE: 2001-08-10
 PRIOR APPLICATION NUMBER: PCT/US00/05883
 PRIOR FILING DATE: 2000-03-08
 PRIOR APPLICATION NUMBER: 60/124,270
PRIOR FILING DATE: 1999-03-12
 NUMBER OF SEQ ID NOS: 1556
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Search completed: January 29, 2004, 02:51:24

Job time : 1088 secs

GenCore version 5.1.6 Copyright (c) 1993 - 2004 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 28, 2004, 20:27:00; Search time 6858 Seconds

(without alignments)

12290.452 Million cell updates/sec

Title: US-10-056-884A-1

Perfect score: 3468

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Gapop 10.0 , Gapext 1.0

Searched: 22781392 segs, 12152238056 residues

Total number of hits satisfying chosen parameters: 45562784

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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3: em_estin:*

4: em_estmu:*

5: em estov:*

6: em estpl:*

7: em estro:*

8: em htc:*

9: gb est1:*

10: gb_est2:*

11: gb htc:*

12: gb_est3:*

13: gb est4:*

14: qb est5:*

15: em estfun:*

16: em_estom:*

17: em_gss_hum:*

18: em_gss_inv:* 19: em_gss_pln:*

20: em_gss_vrt:*

21: em gss fun:* 22: em_gss_mam:*

23: em gss mus:*

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27: em gss vrl:*

28: gb_gss1:* 29: gb_gss2:*

ક

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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С	3	400.2	11.5	592	28	AQ525390	AQ525390 HS 5228 B
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ALIGNMENTS

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            , genomic survey sequence.
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VERSION
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            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
              (bases 1 to 489)
 AUTHORS
            Zhao, S., Adams, M.D., Nierman, W., Malek, J., de Jong, P. and Venter
            ,J.C.
  TITLE
            Use of BAC End Sequences from Library RPCI-11 for Sequence-Ready
            Map Building
  JOURNAL
            Unpublished
COMMENT
            Other GSSs: RPCI-11-318B21.TV
            Contact: Shaying Zhao, William Nierman, Mark Adams
            Department of Eukaryotic Genomics
            The Institute for Genomic Research
            9712 Medical Center Dr., Rockville, MD 20850
            Tel: 301 838 0200
            Fax: 301 838 0208
            Email: hbe@tigr.org
            Clones are derived from the human BAC library RPCI-11. For BAC
            library availability, please contact Pieter de Jong
            (pieter@dejong.med.buffalo.edu). Clones may be purchased from
            BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from
            Research Genet cs (info@resgen.com). BAC end search page:
            http://www.tigr.org/tdb/humgen/bac end search/bac end search.html.
            Seg primer: SP6
            Class: BAC ends.
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REFERENCE AUTHORS TITLE JOURNAL MEDLINE PUBMED	1 C H M 9		

```
REFERENCE
            Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K.,
 AUTHORS
            Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
  TITLE
            Normalization and subtraction of cap-trapper-selected cDNAs to
            prepare full-length cDNA libraries for rapid discovery of new genes
            Genome Res. 10 (10), 1617-1630 (2000)
  JOURNAL
 MEDLINE
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REFERENCE
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 AUTHORS
            Shibata, K., Itoh, M., Aizawa, K., Nagaoka, S., Sasaki, N., Carninci, P.,
            Konno, H., Akiyama, J., Nishi, K., Kitsunai, T., Tashiro, H., Itoh, M.,
            Sumi, N., Ishii, Y., Nakamura, S., Hazama, M., Nishine, T., Harada, A.,
            Yamamoto, R., Matsumoto, H., Sakaguchi, S., Ikegami, T., Kashiwagi, K.,
            Fujiwake, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watahiki, M.,
            Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsuura, S., Kawai, J.,
            Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y.
            RIKEN integrated sequence analysis (RISA) system--384-format
 TITLE
            sequencing pipeline with 384 multicapillary sequencer
  JOURNAL
            Genome Res. 10 (11), 1757-1771 (2000)
            20530913
 MEDLINE
  PUBMED
            11076861
REFERENCE
 AUTHORS
            Kawai, J., Shinagawa, A., Shibata, K., Yoshino, M., Itoh, M., Ishii, Y.,
            Arakawa, T., Hara, A., Fukunishi, Y., Konno, H., Adachi, J., Fukuda, S.,
            Aizawa, K., Izawa, M., Nishi, K., Kiyosawa, H., Kondo, S., Yamanaka, I.,
            Saito, T., Okazaki, Y., Gojobori, T., Bono, H., Kasukawa, T., Saito, R.,
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            Kuehl, P., Lewis, S., Matsuo, Y., Nikaido, I., Pesole, G.,
            Quackenbush, J., Schriml, L.M., Staubli, F., Suzuki, R., Tomita, M.,
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            Baldarelli, R., Barsh, G., Blake, J., Boffelli, D., Bojunga, N.,
            Carninci, P., de Bonaldo, M.F., Brownstein, M.J., Bult, C.,
            Fletcher, C., Fujita, M., Gariboldi, M., Gustincich, S., Hill, D.,
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            Marchionni, L., Mashima, J., Mazzarelli, J., Mombaerts, P., Nordone, P.,
            Ring, B., Ringwald, M., Rodriguez, I., Sakamoto, N., Sasaki, H.,
            Sato, K., Schonbach, C., Seya, T., Shibata, Y., Storch, K.F., Suzuki, H.,
            Toyo-oka, K., Wang, K.H., Weitz, C., Whittaker, C., Wilming, L.,
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            and Hayashizaki, Y.
  TITLE
            Functional annotation of a full-length mouse cDNA collection
  JOURNAL
            Nature 409 (6821), 685-690 (2001)
 MEDLINE
            21085660
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REFERENCE
 AUTHORS
            The FANTOM Consortium and the RIKEN Genome Exploration Research
            Group Phase I & II Team.
            Analysis of the mouse transcriptome based on functional annotation
  TITLE
            of 60,770 full-length cDNAs
  JOURNAL
            Nature 420, 563-573 (2002)
REFERENCE
                (bases 1 to 810)
            Adachi, J., Aizawa, K., Akahira, S., Akimura, T., Arai, A., Aono, H.,
 AUTHORS
            Arakawa, T., Bono, H., Carninci, P., Fukuda, S., Fukunishi, Y.,
            Furuno, M., Hanagaki, T., Hara, A., Hayatsu, N., Hiramoto, K.,
            Hiraoka, T., Hori, F., Imotani, K., Ishii, Y., Itoh, M., Izawa, M.,
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```
Nomura, K., Numazaki, R., Ohno, M., Okazaki, Y., Okido, T., Owa, C.,
           Saito, H., Saito, R., Sakai, C., Sakai, K., Sano, H., Sasaki, D.,
           Shibata, K., Shibata, Y., Shinagawa, A., Shiraki, T., Sogabe, Y.,
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           Yoshino, M., Muramatsu, M. and Hayashizaki, Y.
  TITLE
           Direct Submission
           Submitted (10-JUL-2000) Yoshihide Hayashizaki, The Institute of
  JOURNAL
           Physical and Chemical Research (RIKEN), Laboratory for Genome
           Exploration Research Group, RIKEN Genomic Sciences Center (GSC),
           RIKEN Yokohama Institute; 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama,
           Kanagawa 230-0045, Japan (E-mail:genome-res@gsc.riken.go.jp,
           URL: http://genome.gsc.riken.go.jp/, Tel:81-45-503-9222,
           Fax:81-45-503-9216)
           Please visit our web site (http://genome.gsc.riken.go.jp/) for
COMMENT
           further details.
           cDNA library was prepared and sequenced in Mouse Genome
           Encyclopedia Project of Genome Exploration Research Group in Riken
           Genomic Sciences Center and Genome Science Laboratory in RIKEN.
           Division of Experimental Animal Research in Riken contributed to
           prepare mouse tissues. First strand cDNA was primed with a primer
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           strand cDNA was prepared with the primer adapter of sequence [5'
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           bulk excision from Lambda FLC I. Cloning sites, 5' end: SalI; 3'
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Koya, S., Kurihara, C., Matsuyama, T., Miyazaki, A., Nishi, K.,

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Mahairas, G.G., Wallace, J.C., Smith, K., Swartzell, S., Holzman, T.,
 AUTHORS
           Keller, A., Shaker, R., Furlong, J., Young, J., Zhao, S., Adams, M.D. and
           Hood, L.
 TITLE
           Sequence-tagged connectors: A sequence approach to mapping and
           scanning the human genome
           Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
 JOURNAL
 MEDLINE
           99380589
  PUBMED
           10449764
           Contact: Mahairas GG, Wallace JC, Hood L
COMMENT
           High Throughput Sequencing Center
           University of Washington
           401 Queen Anne Avenue North, Seattle, WA 98109, USA
           Tel: (206) 616-3618
           Fax: (206) 616-3887
           Email: jwallace@u.washington.edu
           Clones are derived from the human BAC library RPCI-11. For BAC
           library availability, please contact Pieter de Jong
           (pieter@dejong.med.buffalo.edu). Clones may be purchased from
           BACPAC Resources (http://bacpac.med.buffalo.edu/ordering bac.htm)
           or from Resear h Genetics (info@resgen.com). BAC end Web Server:
           http://www.htsc.washington.edu
           Plate: 804 row: F column: 10
           Seq primer: T7
           Class: BAC ends
           High quality sequence stop: 592.
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                   /note="Vector: pBACe3.6; Site 1: EcoRI; Site 2: EcoRI;
                   Male blood DNA was isolated from one randomly chosen donor
                   and partially digested with a combination of EcoRI and
                   EcoRI Methylase. Size selected DNA was cloned into the
                   pBACe3.6 vector at EcoRI sites"
BASE COUNT
               157 a
                       139 с
                               133 g
                                        158 t
                                                  5 others
ORIGIN
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                               Score 400.2; DB 28;
 Query Match
                                                    Length 592;
 Best Local Similarity
                        96.2%;
                               Pred. No. 7.6e-34;
 Matches 408; Conservative
                              0; Mismatches
                                             16;
                                                   Indels
                                                             0; Gaps
Qу
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         988 TGCCGACCGCAAGTGGGGTTTCATTACTGTGGGTTACAGAGGATCCTGCACCTTGGGCAG 1047
Qу
             Db
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Qу
             305 GCCAGATAGATACACCTCCAGATTTTATCTCAAATTCAAGCACCTGGAAAGGGCTTTTGA 246
Db
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Qу
            Db
        245 TATGTTGTCAGAGTGTGGATTCCACATGGTGGCCTGTAACTCATCGGTGACAGCATCTTT 186
       1288 CATCAACCAATATACAGATGACAAGATCTGGTCAAGCTACACTGAATATGTCTTCTACCG 1347
Qу
            Db
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       1348 TGAG 1351
Qy
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Db
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CA463745
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          CA463745
                                784 bp
                                        mRNA
                                                linear
                                                       EST 12-NOV-2002
DEFINITION
          AGENCOURT 10724816 NIH MGC 169 Mus musculus cDNA clone
          IMAGE: 6771233 5', mRNA sequence.
ACCESSION
          CA463745
VERSION
          CA463745.1 GI:24920097
KEYWORDS
SOURCE
          Mus musculus (house mouse)
          Mus musculus
 ORGANISM
          Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
          Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
             (bases 1 to 784)
          NIH-MGC http://mgc.nci.nih.gov/.
 AUTHORS
          National Institutes of Health, Mammalian Gene Collection (MGC)
 TITLE
 JOURNAL
          Unpublished
COMMENT
          Contact: Robert Strausberg, Ph.D.
          Email: cgapbs-r@mail.nih.gov
          Tissue Procurement: Dr. Jonathan Kuo, NIMH
           cDNA Library Preparation: Michael Brownstein Laboratory
           cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
           DNA Sequencing by: Agencourt Bioscience Corporation
           Clone distribution: MGC clone distribution information can be
          found through the I.M.A.G.E. Consortium/LLNL at:
          http://image.llnl.gov
          Plate: LLCM3090 row: h column: 16
          High quality sequence stop: 456.
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                  /db xref="taxon:10090"
                  /clone="IMAGE: 6771233"
                  /lab host="DH10B (T1-phage-resistant)"
                  /clone lib="NIH MGC 169"
                  /note="Organ: Testicles; Vector: pDNR-LIB; Site_1: SfiI
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(ggccattatggcc); Site_2: SfiI (ggccgcctcggcc); cDNA made by oligo-dT priming and directionally cloned. 5' and 3' adaptors were used in cloning as follows: 5'-AAGCAGTGGTATCAACGCAGAGTGGCCATTACGGCCGGG-3' and 5'-ATTCTAGAGGCCGAGGCGGCCGACATG-dT(30)NN-3'. Full-length enriched library was constructed using the Clontech Creator SMART kit and size-selected to contain the 0.5 kb size fraction. Library created in thelaboratory of M. Brownstein (NIMH, NIH). Note: this is a NIH MGC Library."

BASE COUNT 244 a 239 c

175 g 121 t 5 others

ORIGIN

Query Match 10.2%; Score 352.2; DB 14; Length 784; Best Local Similarity 85.9%; Pred. No. 8.5e-29; Matches 403; Conservative 0; Mismatches 63; Indels 3; Gaps 1; 1347 GTGAGCCTTCCAGATGGTCACCCTCACACTGCGATTGCTGCTGCAAGAATGGCAAAGGTG 1406 Qу Db 104 GTGAGCCTTCCCGGTGGTCCTCCTCTCATTGTGACTGCTGCTGCAAGAATGGCAAGGGAG 163 Qν 1407 ACAAAGAAGGGGAGAGCGCACGTCTTGCAATGACCTCTCCACATCTAGCTGCGACAGCC 1466 Db 164 ACA---AAGGAGAGAGCGCACCTCCTGCAATGACCTGTCCACTTCCAGCTGTGACAGCC 220 Qу 1467 AGTCTGAGGCCAGCTCTCCCCAGGAGACGGTCATCTGTGGTCCCGTGACACGCCAGACCA 1526 221 AGTCAGAGGCCAGCTCTCCGCAGGAGACGGTGATCTGTGGGCCTGTAACGCGCCAGAGCA 280 Db Qу 1527 ACATCCAGACTCTGGACCGTCCCATCAAGAAGGGCCCTGTCCAGCTGATCCAACAGTCAG 1586 281 ACATCCAGACTCTGGATCGGCCCATCAAGAAAGGTCCGGTGCAGCTGATCCAACAGTCAG 340 Dh 1587 AGATGCGGCGGAAAAGCGACTTACTCCGGATTCTGACTTCAGGCTCCAGGGAATCGAACA 1646 Qy 341 AGATGAGGCGGAAAAGTGACCTGCTCCGGACTCTGACGTCAGGCTCCAGGGAGTCGAACA 400 Db 1647 TGAGCAGCAAAAAAAAGCTGTTAAAGAAAAGCTCTCAATTGAGGAGGAGCTGGAGAAAT 1706 Qy 401 TAAGCAGCAAAAAGAAAGCTGCGAAGGAAAAGCTCTCCATCGAGGAAGAGCTGGAGAAAT 460 Dh 1707 GTATCCAGGATTTCCTAAAAAAAAAAATTCCAGATCGGTTTCCTGAGAGAAAACATCCTT 1766 Qу 461 GTATCCAGGATTTCTTGAAGATAAAAATTCCAGATCGCTTCCCTGAGCGAAAACATCCTT 520 Db Qу 521 GGCAGTCTGAACTTTTACGGGAGTATCATCTATAGGGGGGAGGCTGTGG 569 Db

RESULT 5 BU961910

LOCUS BU961910 778 bp mRNA linear EST 21-OCT-2002 DEFINITION AGENCOURT 10617166 NIH MGC 169 Mus musculus cDNA clone

IMAGE: 6742567 5', mRNA sequence.

ACCESSION BU961910

VERSION BU961910.1 GI:24191482

KEYWORDS EST.

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SOURCE
           Mus musculus (house mouse)
 ORGANISM Mus musculus
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
              (bases 1 to 778)
REFERENCE
           NIH-MGC http://mgc.nci.nih.gov/.
 AUTHORS
           National Institutes of Health, Mammalian Gene Collection (MGC)
 TITLE
 JOURNAL
           Unpublished
COMMENT
           Contact: Robert Strausberg, Ph.D.
           Email: cgapbs-r@mail.nih.gov
           Tissue Procurement: Dr. Jonathan Kuo, NIMH
            cDNA Library Preparation: Michael Brownstein Laboratory
            cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
            DNA Sequencing by: Agencourt Bioscience Corporation
            Clone distribution: MGC clone distribution information can be
           found through the I.M.A.G.E. Consortium/LLNL at:
           http://image.llnl.gov
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                   /clone lib="NIH MGC 169"
                   /note="Organ: Testicles; Vector: pDNR-LIB; Site 1: SfiI
                   (ggccattatggcc); Site 2: SfiI (ggccgcctcggcc); cDNA made
                   by oligo-dT priming and directionally cloned. 5' and 3'
                   adaptors were used in cloning as follows:
                   5'-AAGCAGTGGTATCAACGCAGAGTGGCCATTACGGCCGGG-3' and
                   5'-ATTCTAGAGGCCGAGGCGGCCGACATG-dT(30)NN-3'. Full-length
                   enriched library was constructed using the Clontech
                   Creator SMART kit and size-selected to contain the 0.5 kb
                   size fraction. Library created in thelaboratory of M.
                   Brownstein (NIMH, NIH). Note: this is a NIH MGC Library."
BASE COUNT
              226 a
                       180 c
                               199 g
                                        160 t
                                                 13 others
ORIGIN
 Query Match
                        10.1%;
                               Score 352; DB 13; Length 778;
 Best Local Similarity
                        85.6%; Pred. No. 8.9e-29;
 Matches 404; Conservative
                              0; Mismatches
                                              65;
                                                   Indels
                                                             3; Gaps
                                                                        1;
        1347 GTGAGCCTTCCAGATGGTCACCCTCACACTGCGATTGCTGCTGCAAGAATGGCAAAGGTG 1406
Qy
             Db
         183 GTGAGCCTTCCCGGTGGTCCTCCTCTCATTGTGACTGCTGCCAAGAATGGCAAGGGAG 242
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                  Db
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        1467 AGTCTGAGGCCAGCTCTCCCCAGGAGACGGTCATCTGTGGTCCCGTGACACGCCAGACCA 1526
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Qу
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Qу	1647	TGAGCAGCAAAAAAAAAGCTGTTAAAGAAAAGCTCTCAATTGAGGAGGAGCTGGAGAAAT 17	06
Db	480	TAAGCAGCAAAAAGAAAGCTGCGAAGGAAAAGCTCTCCATCGAGGAAGAGCTGGAGAAAT 53	9
QУ	1707	GTATCCAGGATTTCCTAAAAAAAAATTCCAGATCGGTTTCCTGAGAGAAACATCCTT 17	66
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QУ	1767	GGCAATCTGAACTTTTAAGGAAGTATCATCTATAAGGGAGGG	
Db	600		

RESULT 6 BY714867

LOCUS BY714867 952 bp mRNA linear EST 17-DEC-2002 DEFINITION BY714867 RIKEN full-length enriched, adult male testis Mus musculus cDNA clone 4930434H12 5', mRNA sequence.

ACCESSION BY714867

VERSION BY714867.1 GI:27127984

KEYWORDS EST.

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 952)

AUTHORS Okazaki, Y., Furuno, M., Kasukawa, T., Adachi, J., Bono, H., Kondo, S., Nikaido, I., Osato, N., Saito, R., Suzuki, H., Yamanaka, I., Kiyosawa, H., Yagi, K., Tomaru, Y., Hasegawa, Y., Nogami, A., Schonbach, C., Gojobori, T., Baldarelli, R., Hill, D.P., Bult, C., Hume, D.A., Quackenbush, J., Schriml, L.M., Kanapin, A., Matsuda, H., Batalov, S., Beisel, K.W., Blake, J.A., Bradt, D., Brusic, V., Chothia, C., Corbani, L.E., Cousins, S., Dalla, E., Dragani, T.A., Fletcher, C.F., Forrest, A., Frazer, K.S., Gaasterland, T., Gariboldi, M., Gissi, C., Godzik, A., Gough, J., Grimmond, S., Gustincich, S., Hirokawa, N., Jackson, I.J., Jarvis, E.D., Kanai, A., Kawaji, H., Kawasawa, Y., Kedzierski, R.M., King, B.L., Konagaya, A., Kurochkin, I.V., Lee, Y., Lenhard, B., Lyons, P.A., Maglott, D.R., Maltais, L., Marchionni, L., McKenzie, L., Miki, H., Nagashima, T., Numata, K., Okido, T., Pavan, W.J., Pertea, G.,

Pesole, G., Petrovsky, N., Pillai, R., Pontius, J.U., Qi, D.,

,B.Z., Ringwald, M., Sandelin, A., Schneider, C., Semple, C.A., Setou, M., Shimada, K., Sultana, R., Takenaka, Y., Taylor, M.S., Teasdale, R.D., Tomita, M., Verardo, R., Wagner, L., Wahlestedt, C., Wang, Y., Watanabe, Y., Wells, C., Wilming, L.G., Wynshaw-Boris, A., Yanagisawa, M., Yang, I., Yang, L., Yuan, Z., Zavolan, M., Zhu, Y., Zimmer, A., Carninci, P., Hayatsu, N., Hirozane-Kishikawa, T., Konno, H., Nakamura, M., Sakazume, N., Sato, K., Shiraki, T., Waki, K., Kawai, J., Aizawa, K., Arakawa, T., Fukuda, S., Hara, A., Hashizume, W., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Miyazaki, A., Sakai, K., Sasaki, D., Shibata

Ramachandran, S., Ravasi, T., Reed, J.C., Reed, D.J., Reid, J., Ring

,K., Shinaqawa, A., Yasunishi, A., Yoshino, M., Waterston, R., Lander , E.S., Rogers, J., Birney, E. and Hayashizaki, Y. Analysis of the mouse transcriptome based on functional annotation of 60,770 full-length cDNAs Nature 420, 563-573 (2002) JOURNAL MEDLINE 22354683 PUBMED 12466851 Contact: Yoshihide Hayashizaki Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center(GSC), Yokohama Institute The Institute of Physical and Chemical Research (RIKEN) 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan Tel: 81-45-503-9222 Fax: 81-45-503-9216 Email: genome-res@gsc.riken.go.jp, URL:http://genome.gsc.riken.go.jp/ Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Carninci, P., Fukuda ,S., Hashizume, W., Hayashida, K., Hirozane, T., Hori, F., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Kawai, J., Kojima, Y., Kondo, S., Konno ,H., Koya,S., Miyazaki,A., Murata,M., Nakamura,M., Nomura,K., Numazaki, R., Ohno, M., Ohsato, N., Saito, R., Sakazume, N., Sano, H., Sasaki, D., Sato, K., Shibata, K., Shiraki, T., Tagami, M., Takeda, Y., Waki, K., Watahiki, A., Muramatsu, M. and Hayashizaki, Y. Direct Submission Computational Analysis of Full-Length Mouse cDNAs Compared with Human Genome Sequences Mamm. Genome. 12, 673-677 (2001) Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. Genome Res. 10 (10), 1617-1630 (2000) RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1757-1771 (2000) Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001) cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. Please visit our web site (http://genome.gsc.riken.go.jp) for further details. **FEATURES** Location/Qualifiers source 1. .952 /organism="Mus musculus" /mol type="mRNA" /strain="C57BL/6J" /db xref="taxon:10090" /clone="4930434H12" /sex="male" /tissue type="testis" /dev stage="adult" /lab host="SOLR"

/clone lib="RIKEN full-length enriched, adult male testis"

/note="Site_1: XhoI; Site_2: BamHI; cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken

TITLE

COMMENT

Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. 1st strand cDNA was primed with a primer [5'

BASE COUNT 233 a 259 c 226 g 229 t 5 others ORIGIN

Query Match 10.1%; Score 350.6; DB 14; Length 952; Best Local Similarity 75.5%; Pred. No. 1.1e-28; Matches 506; Conservative 0; Mismatches 142; Indels 22; Gaps				
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DEFINITION UI-R-CAl-bcd-a-05-0-UI.sl UI-R-CAl Rattus norvegicus cDNA clone
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ACCESSION
           BF391086
VERSION
           BF391086.1 GI:11375933
KEYWORDS
           EST.
SOURCE
           Rattus norvegicus (Norway rat)
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           Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
REFERENCE
              (bases 1 to 489)
           1
  AUTHORS
           Bonaldo, M.F., Lennon, G. and Soares, M.B.
 TITLE
           Normalization and subtraction: two approaches to facilitate gene
           discovery
  JOURNAL
           Genome Res. 6 (9), 791-806 (1996)
 MEDLINE
           97044477
  PUBMED
           8889548
COMMENT
           Contact: Soares, MB
           Coordinated Laboratory for Computational Genomics
           University of Iowa
           375 Newton Road , 4156 MEBRF, Iowa City, IA 52242, USA
           Tel: 319 335 8250
           Fax: 319 335 9565
           Email: bento-soares@uiowa.edu
           The sequence contained an oligo-dT track that was present in the
           oligonucleotide that was used to prime the synthesis of first
           strand cDNA and therefore this may represent a bonafide poly A
           tail. The sequence tag present in the cDNA between the NotI site
           and the oligo-dT track served to identify it as a clone from the
           normalized testis library cDNA Library Preparation: M.B. Soares Lab
           Clone distribution: clones will be available through Research
           Genetics (www.resgen.com) The following repetitive elements were
           found in this cDNA sequence: 1-35, >POLY A#Simple repeat
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FEATURES
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/note="Vector: pT7T3D-Pac (Pharmacia) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; The UI-R-CA1 library is a subtracted library derived from the following tissues: thalamus, cerebellum, hypothalamus, medulla, pons, midbrain, cerebral cortex, corpus striatum, testis, and hippocampus. For a detailed description of the library from which this clone was derived, please visit our web site at ratest.eng.uiowa.edu. The subtraction has been previously described in (Bonaldo, Lennon and Soares, Genome Research 6:791-806, 1996)
TAG LIB=UI-R-CA1

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TAG_SEQ=ACGCAG"

BASE COUNT 89 a 123 c 111 g 164 t 2 others ORIGIN

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Qy	1593	GGCGGAAAAGCGACTTACTCCGGATTCTGACTTCAGGCTCCAGGGAATCGAACATGAGCA 1652
Db	303	
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RESULT 8 AK043351

LOCUS AK043351 2332 bp mRNA linear HTC 05-DEC-2002 DEFINITION Mus musculus 7 days neonate cerebellum cDNA, RIKEN full-length

enriched library, clone: A730087N02 product: hypothetical protein, full insert sequence. AK043351 ACCESSION AK043351.1 GI:26335652 VERSION KEYWORDS HTC; CAP trapper. Mus musculus (house mouse) SOURCE ORGANISM Mus musculus Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. REFERENCE AUTHORS Carninci, P. and Hayashizaki, Y. TITLE High-efficiency full-length cDNA cloning Meth. Enzymol. 303, 19-44 (1999) JOURNAL MEDLINE 99279253 10349636 PUBMED REFERENCE **AUTHORS** Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y. TITLE Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes JOURNAL Genome Res. 10 (10), 1617-1630 (2000) 20499374 MEDLINE PUBMED 11042159 REFERENCE 3 Shibata, K., Itoh, M., Aizawa, K., Naqaoka, S., Sasaki, N., Carninci, P., AUTHORS Konno, H., Akiyama, J., Nishi, K., Kitsunai, T., Tashiro, H., Itoh, M., Sumi, N., Ishii, Y., Nakamura, S., Hazama, M., Nishine, T., Harada, A., Yamamoto, R., Matsumoto, H., Sakaguchi, S., Ikegami, T., Kashiwagi, K., Fujiwake, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watahiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsuura, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y. RIKEN integrated sequence analysis (RISA) system--384-format TITLE sequencing pipeline with 384 multicapillary sequencer Genome Res. 10 (11), 1757-1771 (2000) JOURNAL MEDLINE 20530913 PUBMED 11076861 REFERENCE **AUTHORS** Kawai, J., Shinagawa, A., Shibata, K., Yoshino, M., Itoh, M., Ishii, Y., Arakawa, T., Hara, A., Fukunishi, Y., Konno, H., Adachi, J., Fukuda, S., Aizawa, K., Izawa, M., Nishi, K., Kiyosawa, H., Kondo, S., Yamanaka, I., Saito, T., Okazaki, Y., Gojobori, T., Bono, H., Kasukawa, T., Saito, R., Kadota, K., Matsuda, H., Ashburner, M., Batalov, S., Casavant, T., Fleischmann, W., Gaasterland, T., Gissi, C., King, B., Kochiwa, H., Kuehl, P., Lewis, S., Matsuo, Y., Nikaido, I., Pesole, G., Quackenbush, J., Schriml, L.M., Staubli, F., Suzuki, R., Tomita, M., Wagner, L., Washio, T., Sakai, K., Okido, T., Furuno, M., Aono, H., Baldarelli, R., Barsh, G., Blake, J., Boffelli, D., Bojunga, N., Carninci, P., de Bonaldo, M.F., Brownstein, M.J., Bult, C., Fletcher, C., Fujita, M., Gariboldi, M., Gustincich, S., Hill, D., Hofmann, M., Hume, D.A., Kamiya, M., Lee, N.H., Lyons, P., Marchionni, L., Mashima, J., Mazzarelli, J., Mombaerts, P., Nordone, P., Ring, B., Ringwald, M., Rodriguez, I., Sakamoto, N., Sasaki, H., Sato, K., Schonbach, C., Seya, T., Shibata, Y., Storch, K.F., Suzuki, H., Toyo-oka, K., Wang, K.H., Weitz, C., Whittaker, C., Wilming, L., Wynshaw-Boris, A., Yoshida, K., Hasegawa, Y., Kawaji, H., Kohtsuki, S. and Hayashizaki, Y. TITLE Functional annotation of a full-length mouse cDNA collection

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JOURNAL
            Nature 409 (6821), 685-690 (2001)
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   PUBMED
REFERENCE
            The FANTOM Consortium and the RIKEN Genome Exploration Research
 AUTHORS
            Group Phase I & II Team.
            Analysis of the mouse transcriptome based on functional annotation
 TITLE
            of 60,770 full-length cDNAs
  JOURNAL
            Nature 420, 563-573 (2002)
REFERENCE
               (bases 1 to 2332)
            Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Bono, H., Carninci, P.,
 AUTHORS
            Fukuda, S., Furuno, M., Hanagaki, T., Hara, A., Hashizume, W.,
            Hayashida, K., Hayatsu, N., Hiramoto, K., Hiraoka, T., Hirozane, T.,
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            Muramatsu, M. and Hayashizaki, Y.
  TITLE
            Direct Submission
            Submitted (16-JUL-2001) Yoshihide Hayashizaki, The Institute of
  JOURNAL
            Physical and Chemical Research (RIKEN), Laboratory for Genome
            Exploration Research Group, RIKEN Genomic Sciences Center (GSC),
            RIKEN Yokohama Institute; 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama,
            Kanagawa 230-0045, Japan (E-mail:genome-res@gsc.riken.go.jp,
            URL: http://genome.gsc.riken.go.jp/, Tel:81-45-503-9222,
            Fax:81-45-503-9216)
COMMENT
            cDNA library was prepared and sequenced in Mouse Genome
            Encyclopedia Project of Genome Exploration Research Group in Riken
            Genomic Sciences Center and Genome Science Laboratory in RIKEN.
            Division of Experimental Animal Research in Riken contributed to
            prepare mouse tissues.
            Please visit our web site for further details.
            URL:http://genome.gsc.riken.go.jp/
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RESULT 9 AK047519

LOCUS AK047519 2343 bp mRNA linear HTC 05-DEC-2002 DEFINITION Mus musculus 10 days neonate cerebellum cDNA, RIKEN full-length enriched library, clone:B930082J01 product:hypothetical protein, full insert sequence.

ACCESSION AK047519 VERSION AK047519.1 GI:26092232 HTC; CAP trapper. KEYWORDS SOURCE Mus musculus (house mouse) ORGANISM Mus musculus Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. REFERENCE Carninci, P. and Hayashizaki, Y. AUTHORS TITLE High-efficiency full-length cDNA cloning JOURNAL Meth. Enzymol. 303, 19-44 (1999) MEDLINE 99279253 PUBMED 10349636 REFERENCE **AUTHORS** Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y. TITLE Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes Genome Res. 10 (10), 1617-1630 (2000) **JOURNAL** 20499374 MEDLINE PUBMED 11042159 REFERENCE **AUTHORS** Shibata, K., Itoh, M., Aizawa, K., Nagaoka, S., Sasaki, N., Carninci, P., Konno, H., Akiyama, J., Nishi, K., Kitsunai, T., Tashiro, H., Itoh, M., Sumi, N., Ishii, Y., Nakamura, S., Hazama, M., Nishine, T., Harada, A., Yamamoto, R., Matsumoto, H., Sakaguchi, S., Ikegami, T., Kashiwagi, K., Fujiwake, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watahiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsuura, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y. TITLE RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer Genome Res. 10 (11), 1757-1771 (2000) JOURNAL MEDLINE 20530913 PUBMED 11076861 REFERENCE **AUTHORS** Kawai, J., Shinagawa, A., Shibata, K., Yoshino, M., Itoh, M., Ishii, Y., Arakawa, T., Hara, A., Fukunishi, Y., Konno, H., Adachi, J., Fukuda, S., Aizawa, K., Izawa, M., Nishi, K., Kiyosawa, H., Kondo, S., Yamanaka, I., Saito, T., Okazaki, Y., Gojobori, T., Bono, H., Kasukawa, T., Saito, R., Kadota, K., Matsuda, H., Ashburner, M., Batalov, S., Casavant, T., Fleischmann, W., Gaasterland, T., Gissi, C., King, B., Kochiwa, H., Kuehl, P., Lewis, S., Matsuo, Y., Nikaido, I., Pesole, G., Quackenbush, J., Schriml, L.M., Staubli, F., Suzuki, R., Tomita, M., Wagner, L., Washio, T., Sakai, K., Okido, T., Furuno, M., Aono, H., Baldarelli, R., Barsh, G., Blake, J., Boffelli, D., Bojunga, N., Carninci, P., de Bonaldo, M.F., Brownstein, M.J., Bult, C., Fletcher, C., Fujita, M., Gariboldi, M., Gustincich, S., Hill, D., Hofmann, M., Hume, D.A., Kamiya, M., Lee, N.H., Lyons, P., Marchionni, L., Mashima, J., Mazzarelli, J., Mombaerts, P., Nordone, P., Ring, B., Ringwald, M., Rodriguez, I., Sakamoto, N., Sasaki, H., Sato, K., Schonbach, C., Seya, T., Shibata, Y., Storch, K.F., Suzuki, H., Toyo-oka, K., Wang, K.H., Weitz, C., Whittaker, C., Wilming, L., Wynshaw-Boris, A., Yoshida, K., Hasegawa, Y., Kawaji, H., Kohtsuki, S. and Hayashizaki, Y. Functional annotation of a full-length mouse cDNA collection TITLE Nature 409 (6821), 685-690 (2001) JOURNAL MEDLINE 21085660

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REFERENCE
            The FANTOM Consortium and the RIKEN Genome Exploration Research
 AUTHORS
            Group Phase I & II Team.
 TITLE
            Analysis of the mouse transcriptome based on functional annotation
            of 60,770 full-length cDNAs
  JOURNAL
            Nature 420, 563-573 (2002)
               (bases 1 to 2343)
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 TITLE
            Direct Submission
            Submitted (16-JUL-2001) Yoshihide Hayashizaki, The Institute of
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            Physical and Chemical Research (RIKEN), Laboratory for Genome
            Exploration Research Group, RIKEN Genomic Sciences Center (GSC),
            RIKEN Yokohama Institute; 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama,
            Kanagawa 230-0045, Japan (E-mail:genome-res@gsc.riken.go.jp,
            URL: http://genome.gsc.riken.go.jp/, Tel:81-45-503-9222,
            Fax: 81-45-503-9216)
COMMENT
            cDNA library was prepared and sequenced in Mouse Genome
            Encyclopedia Project of Genome Exploration Research Group in Riken
            Genomic Sciences Center and Genome Science Laboratory in RIKEN.
            Division of Experimental Animal Research in Riken contributed to
            prepare mouse tissues.
            Please visit our web site for further details.
            URL:http://genome.gsc.riken.go.jp/
            URL:http://fantom.gsc.riken.go.jp/.
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ACCESSION
VERSION
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REFERENCE
 AUTHORS
          Carninci, P. and Hayashizaki, Y.
 TITLE
          High-efficiency full-length cDNA cloning
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          Meth. Enzymol. 303, 19-44 (1999)
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Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K.,

AUTHORS

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Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
            Normalization and subtraction of cap-trapper-selected cDNAs to
  TITLE
            prepare full-length cDNA libraries for rapid discovery of new genes
            Genome Res. 10 (10), 1617-1630 (2000)
  JOURNAL
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            Kawai, J., Shinagawa, A., Shibata, K., Yoshino, M., Itoh, M., Ishii, Y.,
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  TITLE
            Functional annotation of a full-length mouse cDNA collection
            Nature 409 (6821), 685-690 (2001)
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  AUTHORS
            The FANTOM Consortium and the RIKEN Genome Exploration Research
            Group Phase I & II Team.
  TITLE
            Analysis of the mouse transcriptome based on functional annotation
            of 60,770 full-length cDNAs
  JOURNAL
            Nature 420, 563-573 (2002)
REFERENCE
                (bases 1 to 2584)
  AUTHORS
            Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Bono, H., Carninci, P.,
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            Direct Submission
 TITLE
  JOURNAL
            Submitted (16-JUL-2001) Yoshihide Hayashizaki, The Institute of
            Physical and Chemical Research (RIKEN), Laboratory for Genome
            Exploration Research Group, RIKEN Genomic Sciences Center (GSC),
            RIKEN Yokohama Institute; 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama,
            Kanagawa 230-0045, Japan (E-mail:genome-res@gsc.riken.go.jp,
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            Fax:81-45-503-9216)
COMMENT
            cDNA library was prepared and sequenced in Mouse Genome
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            Genomic Sciences Center and Genome Science Laboratory in RIKEN.
            Division of Experimental Animal Research in Riken contributed to
            prepare mouse tissues.
            Please visit our web site for further details.
            URL:http://genome.gsc.riken.go.jp/
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REFERENCE AUTHORS TITLE JOURNAL MEDLINE PUBMES REFERENCE AUTHORS	E S L E D E	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. Carninci, P. and Hayashizaki, Y. High-efficiency full-length cDNA cloning Meth. Enzymol. 303, 19-44 (1999) 99279253 10349636 Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.

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Normalization and subtraction of cap-trapper-selected cDNAs to
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            prepare full-length cDNA libraries for rapid discovery of new genes
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  AUTHORS
            The FANTOM Consortium and the RIKEN Genome Exploration Research
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            Analysis of the mouse transcriptome based on functional annotation
  TITLE
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  JOURNAL
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            Muramatsu, M. and Hayashizaki, Y.
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  JOURNAL
            Physical and Chemical Research (RIKEN), Laboratory for Genome
            Exploration Research Group, RIKEN Genomic Sciences Center (GSC),
            RIKEN Yokohama Institute; 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama,
            Kanagawa 230-0045, Japan (E-mail:genome-res@gsc.riken.go.jp,
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COMMENT
            cDNA library was prepared and sequenced in Mouse Genome
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            Genomic Sciences Center and Genome Science Laboratory in RIKEN.
            Division of Experimental Animal Research in Riken contributed to
            prepare mouse tissues.
            Please visit our web site for further details.
            URL:http://genome.gsc.riken.go.jp/
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DEFINITION A S ACCESSION B VERSION B KEYWORDS E SOURCE M ORGANISM M REFERENCE 1 AUTHORS N TITLE N JOURNAL U COMMENT C	20713664 973 bp mRNA linear EST 16-JUL-2002 MGENCOURT_8480138 NIH_MGC_129 Mus musculus cDNA clone IMAGE:6310836 b', mRNA sequence. 20713664 30713664.1 GI:21852563 35T. 20713664.1 GI:21852563 20713664.1 GI:21852563 35T. 20713664.1 GI:21852563 35T. 2071366

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	Gi Ke Me	L.M., Fitzhugh, W.M., Fritchman, J.L., Geoghagen, N.S., Glodek, A., nehm, C.L., Hanna, M.C., Hedblom, E., Hinkle, P.S.Jr., Kelley, J.M., elley, J.C., Liu, LI., Marmaros, S.M., Merrick, J.M., oreno-Palanques, R.F., McDonald, L.A., Nguyen, D.T., Pelligrino, S.M., hillips, C.A., Ryder, S.E., Scott, J.L., Saudek, D.M., Shirley, R.,

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Small, K.V., Spriggs, T.A., Utterback, T.R., Weidman, J.F., Li, Y.,
          Bednarik, D.P., Cao, L., Cepeda, M.A., Coleman, T.A., Collins, E.J.,
          Dimke, D., Feng, D.-F., Ferrie, A., Fischer, C., Hastings, G.A., He, W.W.
          , Hu, J.S., Greene, J.M., Gruber, J., Hudson, P., Kim, A.K., Kozak, D.L.,
          Kunsch, C., Hungjun, J., Li, H., Meissner, P.S., Olsen, H., Raymond, L.,
          Wei, Y.F., Wing, J., Xu, C., Yu, G.L., Ruben, S.M., Dillion, P.J., Fannon
          ,M.R., Rosen,C.A., Haseltine,W.A., Fields,C., Fraser,C.M. and
          Venter, J.C.
 TITLE
          Initial assessment of human gene diversity and expression patterns
          based upon 83 million nucleotides of cDNA sequence
 JOURNAL
          Nature 377 (6547 Suppl), 3-174 (1995)
          96026280
 MEDLINE
          7566098
  PUBMED
COMMENT
          Contact: Kerlavage, AR
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          The Institute for Genomic Research
          9712 Medical Center Drive, Rockville, MD 20850 USA
          Tel: 3018699056
          Fax: 3018699423
          Email: arkerlav@tigr.org
          For clone availability, additional sequence and expression
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RESULT 14 BY706433

LOCUS BY706433 424 bp mRNA linear EST 16-DEC-2002 DEFINITION BY706433 RIKEN full-length enriched, adult male testis Mus musculus cDNA clone 1700026A08 5', mRNA sequence.

ACCESSION BY706433

VERSION BY706433.1 GI:27117598

KEYWORDS EST.

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 424)

AUTHORS

Okazaki,Y., Furuno,M., Kasukawa,T., Adachi,J., Bono,H., Kondo,S.,
Nikaido,I., Osato,N., Saito,R., Suzuki,H., Yamanaka,I., Kiyosawa,H.,
Yagi,K., Tomaru,Y., Hasegawa,Y., Nogami,A., Schonbach,C.,
Gojobori,T., Baldarelli,R., Hill,D.P., Bult,C., Hume,D.A.,
Quackenbush,J., Schriml,L.M., Kanapin,A., Matsuda,H., Batalov,S.,
Beisel,K.W., Blake,J.A., Bradt,D., Brusic,V., Chothia,C., Corbani,L.E., Cousins,S., Dalla,E., Dragani,T.A., Fletcher,C.F., Forrest

L.E., Cousins, S., Dalla, E., Dragani, T.A., Fletcher, C.F., Forrest, A., Frazer, K.S., Gaasterland, T., Gariboldi, M., Gissi, C., Godzik, A., Gough, J., Grimmond, S., Gustincich, S., Hirokawa, N., Jackson, I.J., Jarvis, E.D., Kanai, A., Kawaji, H., Kawasawa, Y., Kedzierski, R.M., King, B.L., Konagaya, A., Kurochkin, I.V., Lee, Y., Lenhard, B., Lyons, P.A., Maglott, D.R., Maltais, L., Marchionni, L., McKenzie, L., Miki, H., Nagashima, T., Numata, K., Okido, T., Pavan, W.J., Pertea, G., Pesole, G., Petrovsky, N., Pillai, R., Pontius, J.U., Qi, D., Ramachandran, S., Ravasi, T., Reed, J.C., Reed, D.J., Reid, J., Ring, B.Z., Ringwald, M., Sandelin, A., Schneider, C., Semple, C.A., Setou, M., Shimada, K., Sultana, R., Takenaka, Y., Taylor, M.S., Teasdale, R.D., Tomita, M., Verardo, R., Wagner, L., Wahlestedt, C., Wang, Y., Watanabe, Y., Wells, C., Wilming, L.G., Wynshaw-Boris, A., Yanagisawa, M., Yang, I., Yang, L., Yuan, Z., Zavolan, M., Zhu, Y., Zimmer, A.,

,M., Sakazume,N., Sato,K., Shiraki,T., Waki,K., Kawai,J., Aizawa,K., Arakawa,T., Fukuda,S., Hara,A., Hashizume,W., Imotani,K., Ishii,Y., Itoh,M., Kagawa,I., Miyazaki,A., Sakai,K., Sasaki,D., Shibata,K., Shinagawa,A., Yasunishi,A., Yoshino,M., Waterston,R., Lander,E.S., Rogers,J., Birney,E. and Hayashizaki,Y.

Carninci, P., Hayatsu, N., Hirozane-Kishikawa, T., Konno, H., Nakamura

TITLE Analysis of the mouse transcriptome based on functional annotation of 60,770 full-length cDNAs

JOURNAL Nature 420, 563-573 (2002)

MEDLINE 22354683 PUBMED 12466851

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Fax: 81-45-503-9216

Email: genome-res@gsc.riken.go.jp, URL:http://genome.gsc.riken.go.jp/

Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Carninci, P., Fukuda, S., Hashizume, W., Hayashida, K., Hirozane, T., Hori, F., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Kawai, J., Kojima, Y., Kondo, S., Konno, H., Koya, S., Miyazaki, A., Murata, M., Nakamura, M., Nomura, K., Numazaki, R., Ohno, M., Ohsato, N., Saito, R., Sakazume, N., Sano, H., Sasaki, D., Sato, K., Shibata, K., Shiraki, T., Tagami, M., Takeda, Y., Waki, K., Watahiki, A., Muramatsu, M. and Hayashizaki, Y. Direct Submission

Computational Analysis of Full-Length Mouse cDNAs Compared with Human Genome Sequences Mamm. Genome. 12, 673-677 (2001)

Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. Genome Res. 10 (10), 1617-1630 (2000)

RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1757-1771 (2000)

Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)

cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues.

Please visit our web site (http://genome.gsc.riken.go.jp) for further details.

FEATURES

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Location/Qualifiers

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BASE COUNT 133 a 98 c 114 g 79 t

ORIGIN

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RESULT 15 AK006368

Db

LOCUS AK006368 422 bp mRNA linear HTC 05-DEC-2002 DEFINITION Mus musculus adult male testis cDNA, RIKEN full-length enriched library, clone:1700026A08 product:inferred: RIKEN cDNA 4930434H12 gene / putative [Mus musculus], full insert sequence.

ACCESSION AK006368

VERSION AK006368.1 GI:12839431

KEYWORDS HTC; CAP trapper.

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1

AUTHORS Carninci, P. and Hayashizaki, Y.

TITLE High-efficiency full-length cDNA cloning

JOURNAL Meth. Enzymol. 303, 19-44 (1999)

407 TA-ACAACTAACGGTCCAC 424

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99279253
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   PUBMED
REFERENCE
            Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K.,
 AUTHORS
            Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
            Normalization and subtraction of cap-trapper-selected cDNAs to
  TITLE
            prepare full-length cDNA libraries for rapid discovery of new genes
            Genome Res. 10 (10), 1617-1630 (2000)
  JOURNAL
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REFERENCE
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            Shibata, K., Itoh, M., Aizawa, K., Nagaoka, S., Sasaki, N., Carninci, P.,
  AUTHORS
            Konno, H., Akiyama, J., Nishi, K., Kitsunai, T., Tashiro, H., Itoh, M.,
            Sumi, N., Ishii, Y., Nakamura, S., Hazama, M., Nishine, T., Harada, A.,
            Yamamoto, R., Matsumoto, H., Sakaguchi, S., Ikegami, T., Kashiwagi, K.,
            Fujiwake, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watahiki, M.,
            Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsuura, S., Kawai, J.,
            Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y.
  TITLE
            RIKEN integrated sequence analysis (RISA) system--384-format
            sequencing pipeline with 384 multicapillary sequencer
  JOURNAL
            Genome Res. 10 (11), 1757-1771 (2000)
  MEDLINE
            20530913
            11076861
   PUBMED
REFERENCE
            4
  AUTHORS
            Kawai, J., Shinagawa, A., Shibata, K., Yoshino, M., Itoh, M., Ishii, Y.,
            Arakawa, T., Hara, A., Fukunishi, Y., Konno, H., Adachi, J., Fukuda, S.,
            Aizawa, K., Izawa, M., Nishi, K., Kiyosawa, H., Kondo, S., Yamanaka, I.,
            Saito, T., Okazaki, Y., Gojobori, T., Bono, H., Kasukawa, T., Saito, R.,
            Kadota, K., Matsuda, H., Ashburner, M., Batalov, S., Casavant, T.,
            Fleischmann, W., Gaasterland, T., Gissi, C., King, B., Kochiwa, H.,
            Kuehl, P., Lewis, S., Matsuo, Y., Nikaido, I., Pesole, G.,
            Quackenbush, J., Schriml, L.M., Staubli, F., Suzuki, R., Tomita, M.,
            Wagner, L., Washio, T., Sakai, K., Okido, T., Furuno, M., Aono, H.,
            Baldarelli, R., Barsh, G., Blake, J., Boffelli, D., Bojunga, N.,
            Carninci, P., de Bonaldo, M.F., Brownstein, M.J., Bult, C.,
            Fletcher, C., Fujita, M., Gariboldi, M., Gustincich, S., Hill, D.,
            Hofmann, M., Hume, D.A., Kamiya, M., Lee, N.H., Lyons, P.,
            Marchionni, L., Mashima, J., Mazzarelli, J., Mombaerts, P., Nordone, P.,
            Ring, B., Ringwald, M., Rodriguez, I., Sakamoto, N., Sasaki, H.,
             Sato, K., Schonbach, C., Seya, T., Shibata, Y., Storch, K.F., Suzuki, H.,
            Toyo-oka, K., Wang, K.H., Weitz, C., Whittaker, C., Wilming, L.,
            Wynshaw-Boris, A., Yoshida, K., Hasegawa, Y., Kawaji, H., Kohtsuki, S.
             and Hayashizaki, Y.
  TITLE
             Functional annotation of a full-length mouse cDNA collection
  JOURNAL
            Nature 409 (6821), 685-690 (2001)
            21085660
  MEDLINE
   PUBMED
             11217851
REFERENCE
  AUTHORS
            The FANTOM Consortium and the RIKEN Genome Exploration Research
            Group Phase I & II Team.
  TITLE
            Analysis of the mouse transcriptome based on functional annotation
            of 60,770 full-length cDNAs
  JOURNAL
            Nature 420, 563-573 (2002)
REFERENCE
                (bases 1 to 422)
  AUTHORS
            Adachi, J., Aizawa, K., Akahira, S., Akimura, T., Arai, A., Aono, H.,
            Arakawa, T., Bono, H., Carninci, P., Fukuda, S., Fukunishi, Y.,
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Furuno, M., Hanagaki, T., Hara, A., Hayatsu, N., Hiramoto, K.,

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Kasukawa, T., Kato, H., Kawai, J., Kojima, Y., Konno, H., Kouda, M.,
           Koya, S., Kurihara, C., Matsuyama, T., Miyazaki, A., Nishi, K.,
           Nomura, K., Numazaki, R., Ohno, M., Okazaki, Y., Okido, T., Owa, C.,
           Saito, H., Saito, R., Sakai, C., Sakai, K., Sano, H., Sasaki, D.,
           Shibata, K., Shibata, Y., Shinagawa, A., Shiraki, T., Sogabe, Y.,
           Suzuki, H., Tagami, M., Tagawa, A., Takahashi, F., Tanaka, T.,
           Tejima, Y., Toya, T., Yamamura, T., Yasunishi, A., Yoshida, K.,
           Yoshino, M., Muramatsu, M. and Hayashizaki, Y.
 TITLE
           Direct Submission
           Submitted (10-JUL-2000) Yoshihide Hayashizaki, The Institute of
 JOURNAL
           Physical and Chemical Research (RIKEN), Laboratory for Genome
           Exploration Research Group, RIKEN Genomic Sciences Center (GSC),
           RIKEN Yokohama Institute; 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama,
           Kanagawa 230-0045, Japan (E-mail:genome-res@gsc.riken.go.jp,
           URL: http://genome.gsc.riken.go.jp/, Tel:81-45-503-9222,
           Fax:81-45-503-9216)
COMMENT
           Please visit our web site (http://genome.gsc.riken.go.jp/) for
           further details.
           cDNA library was prepared and sequenced in Mouse Genome
           Encyclopedia Project of Genome Exploration Research Group in Riken
           Genomic Sciences Center and Genome Science Laboratory in RIKEN.
           Division of Experimental Animal Research in Riken contributed to
           prepare mouse tissues. First strand cDNA was primed with a primer
           prepared by using trehalose thermo-activated reverse transcriptase
           and subsequently enriched for full-length by cap-trapper. Second
           strand cDNA was prepared with the primer adapter of sequence[5'
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Hiraoka, T., Hori, F., Imotani, K., Ishii, Y., Itoh, M., Izawa, M.,

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